



Cite this: *Org. Biomol. Chem.*, 2018, **16**, 5286

Total synthesis of the proposed structure of talarolide A†

Shengping Zhang,^a Luis M. De Leon Rodriguez,^{id} ^b Renjie Huang,^a Ivanhoe K. H. Leung,^{id} ^a Paul W. R. Harris^{id} ^{a,b,c} and Margaret A. Brimble^{id} ^{*a,b,c}

The proposed structure of talarolide A, a cycloheptapeptide featuring a hydroxamate moiety within the peptide backbone, was successfully synthesized. An initial attempt to synthesize a linear peptide precursor containing a C-terminal *N*-benzyloxy glycine residue was problematic due to an unreported on-resin reduction of *N*-benzyloxy glycine to glycine. After repositioning the peptide cyclization point, a new linear peptide sequence was successfully prepared using Fmoc-solid-phase peptide synthesis. Subsequent solution-phase cyclization and removal of protecting groups furnished the synthetic talarolide A in good yield. Despite the mismatch of the NMR data between the synthetic talarolide A and the natural product, a detailed structural analysis using 2D NMR spectroscopy, together with re-synthesis of the same synthetic material using two additional cyclization sites, confirmed that our synthetic product has the reported structure of talarolide A.

Received 25th May 2018,
Accepted 29th June 2018

DOI: 10.1039/c8ob01230j

rs.c.li/obc

Introduction

The *N*-hydroxylation of peptide backbones is an important strategy for peptide post-translational modification which has mainly been found in the metabolites of microorganisms.^{1–4} Naturally-occurring peptides containing the *N*-hydroxy amide moiety have been reported as potential antibacterial and anti-tumor agents.^{2,3,5,6} Additionally, they can act as siderophores which chelate and transport metal ions essential for cell growth and proliferation.¹ Synthetic peptides containing the *N*-hydroxy amide unit, also exhibit a wide spectrum of bioactivities including inhibition of metalloproteases and HIV protease, as well as immune suppression.^{7–10} It is proposed that the *N*-hydroxy amide functionality not only serves as a strong proton donor participating in hydrogen bonding and metal chelation but also confers enhanced stability to enzymatic degradation compared to its cognate amide counterpart.^{11,12}

Talarolide A (**1**) is a cyclic heptapeptide isolated from an Australian marine tunicate-associated fungus, *Talaromyces* sp. (CMB TU011) (Fig. 1).¹³ Its overall structural elucidation was recently reported by Capon *et al.* using *de novo* spectroscopic analysis and a combination of C₃, C₁₈, and 2D C₃ Marfey's analyses.¹³ The relevant structural features of **1** are the presence of a characteristic *N*-hydroxy amide moiety within the peptide backbone, along with multiple *N*-methyl amino acids. Reports on naturally derived cyclic peptides containing both *N*-hydroxy and *N*-methyl residues are scarce and some of these peptides exhibit potent inhibitory activity against Gram-positive bacteria,² *Mycobacterium tuberculosis* (TB)¹⁴ and the oxytocin receptor.⁴ More importantly, only one recent account of their synthesis has been published to date, which involved the use of corrosive coupling reagents, orthogonal protecting groups

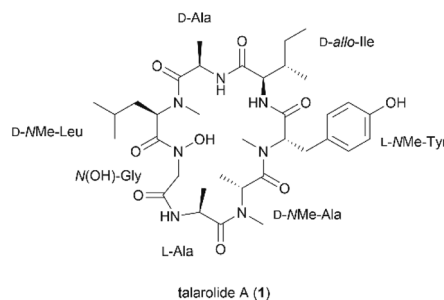


Fig. 1 Structure of talarolide A (**1**).

^aSchool of Chemical Sciences, The University of Auckland, 23 Symonds St, Auckland, 1142, New Zealand. E-mail: m.brimble@auckland.ac.nz; Fax: +649 3737422; Tel: +64 9 3737599

^bMaurice Wilkins Centre for Molecular Biodiscovery, The University of Auckland, Auckland, 1142, New Zealand

^cSchool of Biological Sciences, The University of Auckland, 3A Symonds Street, Auckland 1010, New Zealand

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8ob01230j



and a multi-step synthesis of the *N*-hydroxylated amino acid.¹⁵ We therefore embarked on the first total synthesis of talarolide A (**1**) in order to confirm the proposed structure and establish a robust synthetic route towards this unusual family of peptides.

Results and discussion

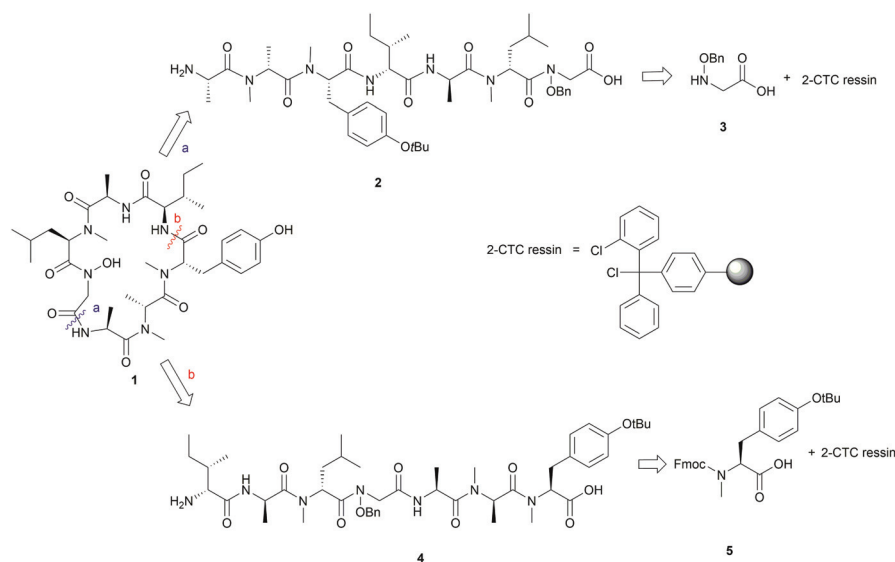
We envisaged that **1** could be constructed employing Fmoc-solid-phase peptide synthesis (SPPS) to access the linear peptide precursor followed by a solution-phase head-to-tail cyclization. Specifically, a side-chain-protected linear peptide could be initially assembled on a hyperacid-labile resin and the *N*-hydroxy amide moiety could be introduced by incorporating an *N*-benzyloxy glycine (**3**) building block into the sequence; after releasing the linear peptide precursor from the resin, **1** can be obtained through a solution-phase macrolactamization followed by subsequent final deprotection (Scheme 1). Moreover, the presence of multiple *N*-methyl amino acids in the peptide sequence could improve the efficiency of the peptide cyclization step by inducing a β -turn conformation that reduces the average distance between the C- and N-terminal residues.¹⁶

Careful selection of the appropriate cyclization point is the key decision in order to successfully synthesize cyclic peptides. In this case, disconnections of **1** at sites that involved *N*-hydroxy and *N*-methyl amide bonds were excluded as it is known that cyclization at those sites is undesirable due to severe C-terminal epimerization and low reaction efficiency caused by the steric hindrance and decreased nucleophilicity of the terminal amine.^{16–19} Among the three remaining cyclization sites, the junction between *N*-OH-Gly and L-Ala was initially chosen, as the derived linear precursor **2** contains an

achiral *N*-OH-Gly residue at the C-terminus, thus reducing the risk of epimerization during peptide cyclization (see disconnection a in Scheme 1).

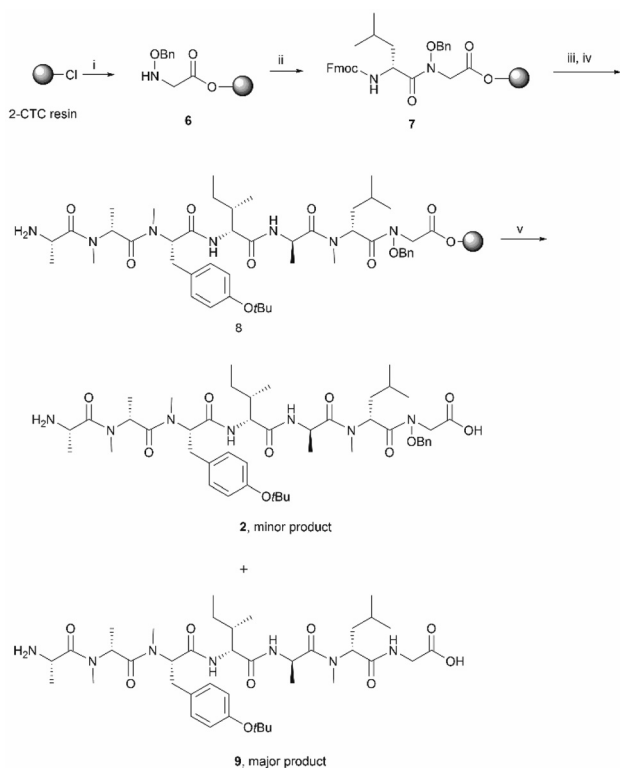
The synthesis of **2** commenced with the attachment of the *N*-benzyloxy glycine (**3**) building block onto a 2-chlorotrityl chloride (2-CTC) resin (Scheme 2). The building block **3** was readily prepared from bromoacetic acid *tert*-butyl ester and *O*-benzylhydroxylamine according to reported methods.^{20,21} The absence of a protecting group on the benzyloxy amine of **3** did not affect the esterification of the first amino acid with 2-CTC resin due to its modified nucleophilicity and steric hindrance compared to unsubstituted amines.²² However, these intrinsic properties of the benzyloxy amine also posed a challenge during the following acylation step. Only strong acylating reagents, such as an acid chloride, a mixed anhydride, or *O*-(7-azabenzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate (HATU), were reported to be successful for the formation of an *N*-hydroxy amide bond.^{10,21,22} In this work, the coupling of Fmoc-D-NMe-Leu to the resin bound benzyloxy glycine was accomplished using HATU as the coupling reagent for 24 h. However, a second coupling was necessary to drive the reaction to completion and thus obtain **7** (Scheme 2). Pleasingly, the long reaction time did not give rise to significant epimerization which has been frequently observed in the acylation of sterically bulky amino acids.^{19,23}

Subsequent peptide elongation with the remaining amino acids was then performed using HATU as the coupling reagent and 20% piperidine in DMF for *N*^α-Fmoc-deprotection, which led to the resin bound peptide **8**. However, LC-MS analysis of a crude sample resulting from the cleavage of **2** from the resin showed an unexpected loss of the benzyloxyl fragment of the *N*-hydroxy glycine residue which resulted in the corresponding C-terminal glycine derivative **9** as the major product (Scheme 2). This side reaction was attributed to the repetitive



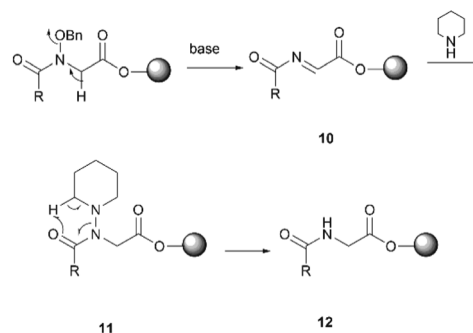
Scheme 1 First (a) and second (b) retrosynthesis of talarolide A (**1**).





Scheme 2 Synthesis of the initial linear peptide precursor **2**. Reagents and conditions: (i) **3** (2 equiv.), DIPEA (5 equiv.), CH_2Cl_2 , RT, 2 h; (ii) Fmoc-NMe-D-Leu (4 equiv.), HATU (4 equiv.), DIPEA (8 equiv.), RT, 2×1 day; (iii) iterative Fmoc-SPPS (a) 20% piperidine in DMF, RT, 2×5 min; (b) Fmoc-AA-OH (4 equiv.), HATU (4 equiv.), DIPEA (8 equiv.), DMF, RT, 1 h; (iv) 20% piperidine in DMF, RT, 2×5 min; (v) HFIP/ CH_2Cl_2 (1 : 4), RT, 1 h.

treatment with piperidine, given that accumulation of the corresponding C-terminal reduced product was observed after each amino acid coupling cycle. For instance, while approximately 50% of the reduced tetrapeptide was observed after cleavage of the NH_2 -D-*allo*-Ile-D-Ala-D-NMe-Leu-(NOBn)Gly-OH peptide from the resin, the reduced peptide **9** (found 776.4 $[M + H]^+$, calcd 776.5) accounted for ~80% of the final product (Fig. S1, S2,† Scheme 2). The loss of the benzyloxy fragment was attributed to the thermal reduction of the *N*-benzyloxy glycine residue to glycine, which has not been reported in previous synthesis of hydroxamate-containing peptides.^{10,21} However, it is important to note that in those previous syntheses the *N*-hydroxy residues were located in the middle of the peptide sequence rather than in the C-terminal region, as is the case presented in Scheme 2. We therefore postulated that the ester bond linkage between the first amino acid and the resin rendered the *N*-acyl-*N*-benzyloxy glycine moiety prone to a base-induced elimination to afford the corresponding α -acylimine intermediate **10**.^{22,24–26} After attack by a “soft nucleophile” such as piperidine, an *N*-substituted glycine derivative **11** is generated, which then undergoes further thermal reduction to give the corresponding glycine derivative **12** (Scheme 3).^{22,27}



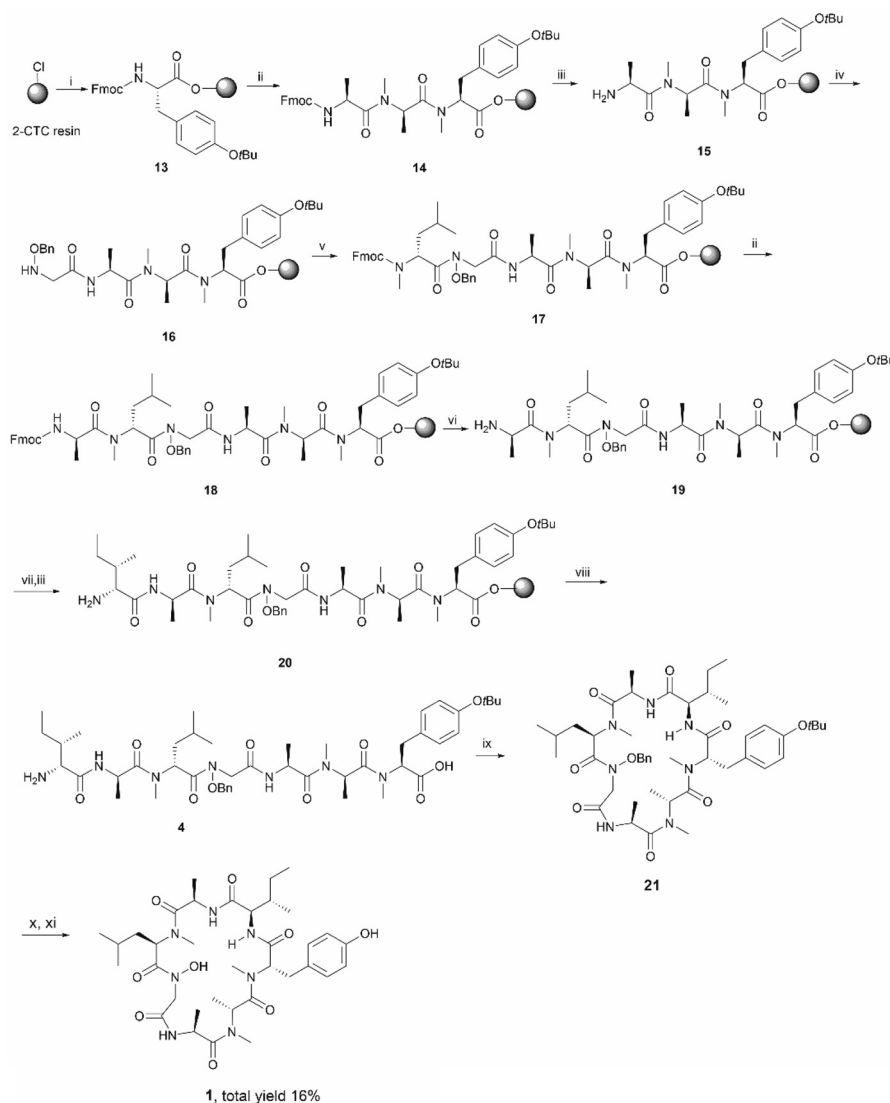
Scheme 3 The proposed mechanism for the reduction of the C-terminal *N*-acyl-*N*-benzyloxy glycine residue.

Having established that the C-terminal *N*-acyl-*N*-benzyloxy glycine ester was not stable to repetitive treatment with bases such as DIPEA and piperidine, we decided to change the cyclization site to the junction between *N*Me-Tyr and *D*-*allo*-Ile so that the building block **3** would be repositioned in the middle of peptide sequence (Scheme 1, path b). The resulting linear precursor **4** was then assembled on a 2-CTC resin as shown in Scheme 4. The coupling of **3** to the resin-bound tripeptide **15** was performed using a mixture of **3**, *N,N'*-diisopropylcarbodiimide (DIC) and 6-chloro-1-hydroxybenzotriazole (6-Cl-HOBT) in DMF for 24 h as powerful coupling reagents, such as HATU, might result in self-condensation of **3** during the long reaction time. For the Fmoc deprotection of **18** we encountered significant diketopiperazine formation which could not be alleviated using milder basic conditions (50% morpholine in DMF and 5% piperazine in DMF containing 0.1 M 6-Cl-HOBT). Fortunately, this undesirable side reaction was minimized using 20% piperidine in DMF and a short deprotection time (2×30 seconds), thereby affording **19** in almost quantitative yield (Scheme 4).

The protected linear precursor **4** (found 882.5 $[M + H]^+$, calcd 882.5) was released from the resin upon treatment with 20% hexafluoroisopropanol (HFIP) in CH_2Cl_2 and the crude peptide was taken to the next step without further purification. The linear peptide **4** was then subjected to peptide cyclization in solution using (benzotriazol-1-yloxy)tripyrrolidinophosphonium hexafluorophosphate (PyBOP) as the cyclizing reagent under high dilution conditions (0.75 mM).^{28,29} LC-MS analysis of the crude reaction mixture indicated the rapid formation of a single product which exhibited the correct mass of the protected cyclic peptide **21**. It is important to note that C-terminal epimerization was not detected during the synthesis, which echoed our hypothesis that a ring-closing conformation would be favored in a peptide sequence encompassing multiple *N*-methyl amino acids.

Subsequent deprotection of the *t*-butyl and benzyl group of **21**, using 90% formic acid aqueous solution and palladium-catalyzed hydrogenation respectively, furnished the desired final product **1** in good yield (16% overall yield based on the determined loading of 2-CTC resin (0.48 mmol g^{-1})) (Scheme 4). It is also worth noting that cyclic peptide **21** was





Scheme 4 The second synthetic route towards talarolide A (**1**). Reagents and conditions: (i) Fmoc-NMe-Tyr(OtBu)-OH (2 equiv.), DIPEA (5 equiv.), CH₂Cl₂, RT, 1 h; (ii) iterative Fmoc-SPPS ((a) 20% piperidine in DMF, RT, 2 × 5 min; (b) Fmoc-AA-OH (4 equiv.), HATU (4 equiv.), DIPEA (8 equiv.), DMF, RT, 1 h); (iii) 20% piperidine in DMF, RT, 2 × 5 min; (iv) **3** (4 equiv.), DIC (4 equiv.), 6-Cl-HOBT (4 equiv.), DMF, RT, 24 h; (v) Fmoc-NMe-D-Leu (4 equiv.), HATU (4 equiv.), DIPEA (8 equiv.), RT, 2 × 1 day; (vi) 20% piperidine in DMF, RT, 2 × 30 s; (vii) Fmoc-D-*allo*-Ile (4 equiv.), HATU (4 equiv.), DIPEA (8 equiv.), DMF, RT, 1 h; (viii) HFIP/CH₂Cl₂ (1 : 4), RT, 1 h; (ix) PyBOP (3 equiv.), DIPEA (5 equiv.), DMF, RT, 1 day; (x) 90% formic acid in water, RT, 40 min; (xi) Pd-C, H₂, MeOH, RT, 90 min.

unstable in typical TFA-mediated *t*-butyl removal conditions, under which conditions it underwent a ring opening reaction to generate the corresponding linear peptide **2**.

The synthetic talarolide A (**1**) was then characterized by ¹H and ¹³C NMR spectroscopy and the obtained data were compared to those reported for the natural product. Unfortunately, the ¹H and ¹³C NMR spectra of **1** did not match those of the natural product. More importantly, two sets of signals with an approximate 1 : 1 ratio were observed in the ¹H-NMR spectrum of **1**, which were attributed to the existence of potential conformers as the *N*-methyl or *N*-hydroxy amide moieties in talarolide A (**1**) are capable of assuming either a *cis* or *trans* conformation (Fig. 2).^{30–32}

In order to support this hypothesis and unambiguously assign all the proton signals in the NMR spectrum to each individual conformer, full NMR characterization of the synthetic talarolide A (**1**) was carried out. The correct peptide sequences were confirmed for both conformers by HMBC. Moreover, the ROESY spectrum of **1** rendered a comprehensive profile of the spatial relationship between all the protons, from which the isomerization state for each *N*-methyl and *N*-hydroxy amide bonds can be deduced. As shown in Fig. 2, a correlation between the H^α of *N*(OH)-Gly and *D*-NMe-Leu residue (**b** and **b'** in Fig. 2) was found in both conformers, thus indicating a *cis* conformation for the *N*-hydroxy amide moiety. In addition, all the amide bonds that involved *N*-methyl substi-



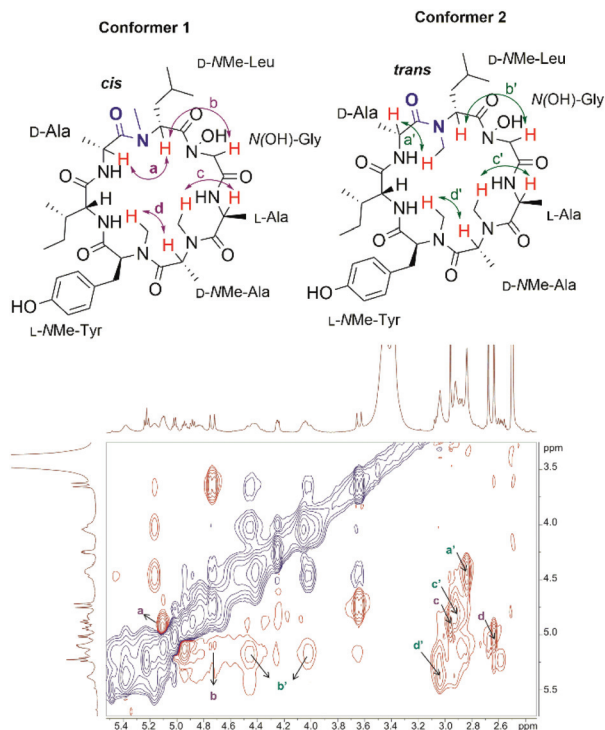


Fig. 2 Expanded view of the ROESY spectrum of synthetic talarolide A (1).

tution assumed the *trans* conformation in both conformers, except the one between the *D*-NMe-Leu and *D*-Ala residues. The correlation observed between the NCH_3 proton of *D*-NMe-Leu and the H^α of *D*-Ala in conformer 2 (a' in Fig. 2) suggested the presence of a *trans* conformation while a *cis* conformation was identified in conformer 1, which was evidenced by the correlation between the H^α of *D*-NMe-Leu and *D*-Ala (a in Fig. 2). This finding pinpointed the position of amide bond where the *cis/trans* isomerization occurred in the synthetic talarolide A (1), thus providing deep insight into the structural difference between the two conformers observed in the NMR experiment. Unfortunately, the published ROESY spectrum of the natural product did not provide sufficient information regarding the isomerization state of each amide bond and only a *trans* conformation was determined for the amide bond between *D*-NMe-Leu and *D*-Ala residue.¹³

Variable temperature NMR experiments were then carried out. As shown in Fig. 3, the proton resonances of these two conformers were well resolved at 27 °C, possibly due to the slow interconversion between the *cis* and *trans* conformations (see below). As the temperature increased, the rate of interconversion increased, leading to peak broadening and signal coalescence, as demonstrated by the ratio between the two conformers, which went from 1 : 1 at RT to 1 : 0.5 at 60 °C (the peaks in the $\text{C}^\alpha\text{-H}$ region were integrated). However, a complete fusion of these two conformers had not been achieved even when the temperature reached 60 °C, thus indicating a relatively high energy barrier between them. A concentration-

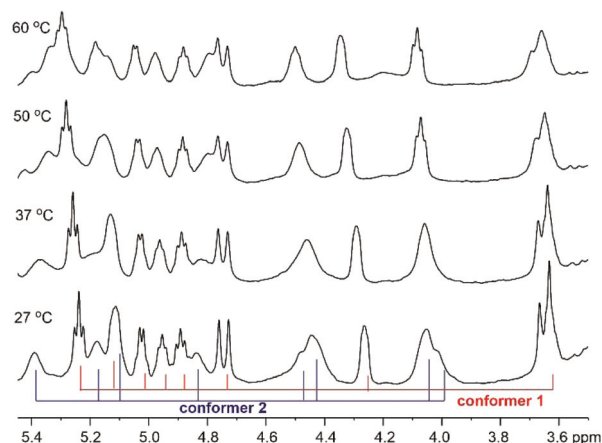
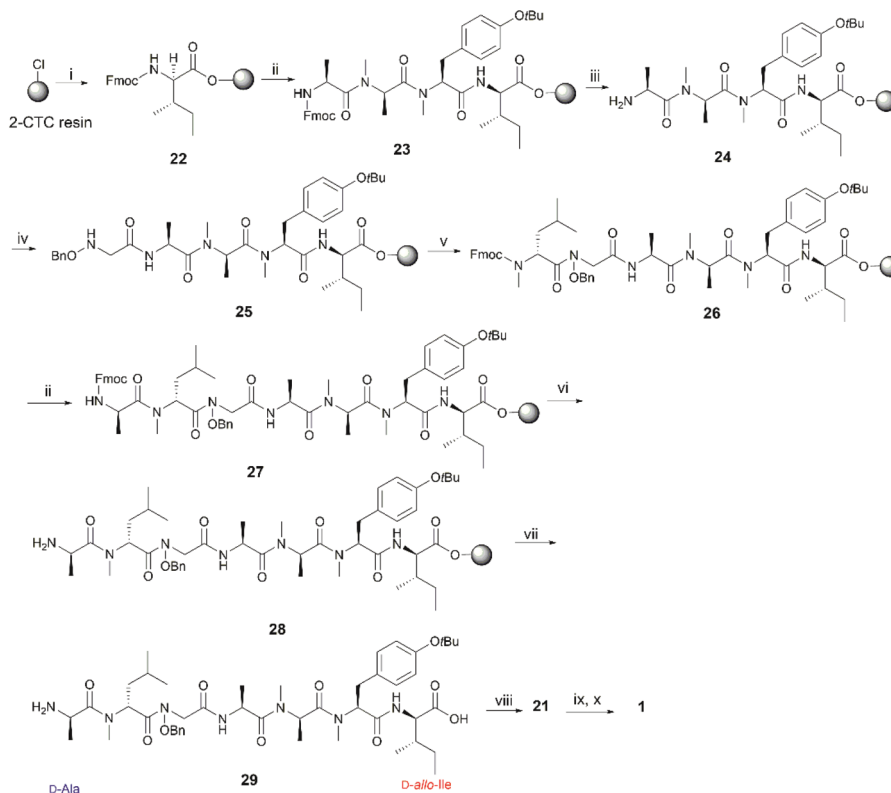


Fig. 3 $\text{C}^\alpha\text{-H}$ region of the ^1H NMR spectra (500 MHz, $\text{DMSO-}d_6$) of synthetic talarolide A (1) at 27 °C, 37 °C, 50 °C and 60 °C.

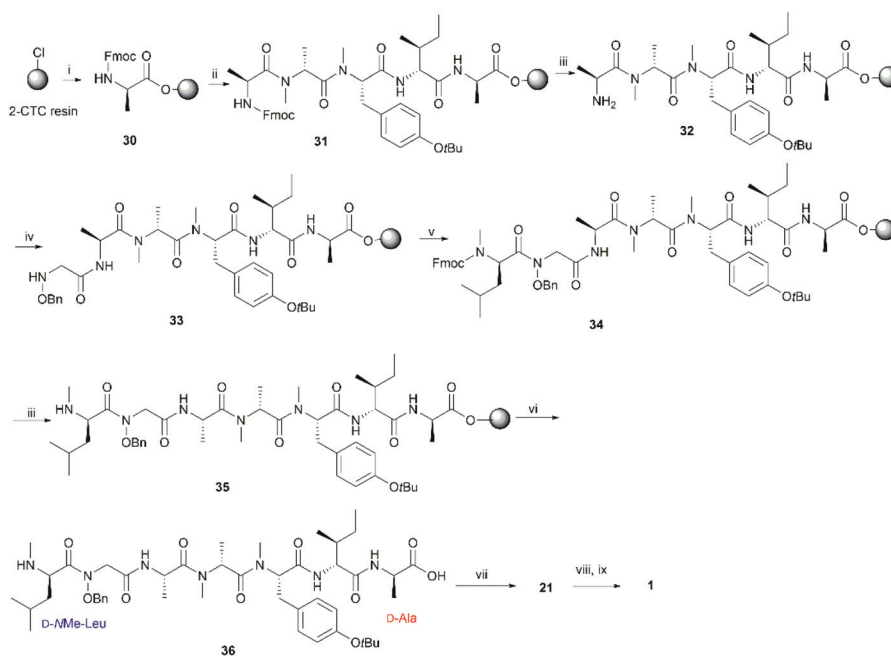
dependent NMR experiment was performed to examine the effect of peptide concentration on its ^1H -NMR spectrum. The obtained ^1H -NMR spectra of 1 at three different concentrations did not exhibit significant variation, hence it is concluded that peptide concentration was not a factor that could explain the differences observed in the NMR spectra of the synthetic and natural talarolide A peptides (Fig. S12†).

In order to further confirm the validity of our synthesis of 1 and to evaluate the impact of different cyclizing sites on peptide conformation, we also synthesized 1 using two different disconnecting points (Schemes 5 and 6). Both linear precursors 29 and 36 were successfully prepared on 2-CTC resin using the same method as described above for the synthesis of 4. Macrolactamization of 29 in DMF (0.75 mM) proceeded smoothly using PyBOP while cyclization of 36 using the same conditions failed to give the desired peptide 21 probably due to the steric hindrance of the *N*-methylated amine at *N*-terminus. Linear peptide 36 was successfully cyclized using HATU, but significant C-terminal epimerization was observed during this process. This phenomenon agreed well with previous reports where severe epimerization took place during the cyclization of peptides that contained an *N*-methyl amino acid at the *N*-terminus.^{17–19} After removal of protecting groups, the final product 1 derived from 29 and 36 was obtained in 7% and 2% overall yield respectively (based on the corresponding loading of 2-CTC resin (0.41 mmol g^{-1} and 0.55 mmol g^{-1})). Finally, the samples of synthetic talarolide A (1) obtained from these two different cyclizing points were characterized by ^1H -NMR, both of which exhibited the same ^1H -NMR spectrum as the one we obtained *via* the synthetic route outlined in Scheme 4, thereby ruling out the existence of possible epimers in the final product (Fig. 4). Moreover, given the fact that the two conformers were consistently observed in all the ^1H -NMR spectra of 1 (Fig. 4), it was concluded that the selection of a different cyclization site for execution of the synthesis had a negligible effect on the conformation of the final peptide.





Scheme 5 Re-synthesis of **1** from the junction between *D*-*allo*-Ile and *D*-Ala. Reagents and conditions: (i) Fmoc-*D*-*allo*-Ile-OH (2 equiv.), DIPEA (5 equiv.), CH₂Cl₂, RT, 1 h; (ii) iterative Fmoc-SPPS ((a) 20% piperidine in DMF, RT, 2 × 5 min; (b) Fmoc-AA-OH (4 equiv.), HATU (4 equiv.), DIPEA (8 equiv.), DMF, RT, 1 h); (iii) 20% piperidine in DMF, RT, 2 × 5 min; (iv) **3** (4 equiv.), DIC (4 equiv.), 6-Cl-HOBt (4 equiv.), DMF, RT, 24 h; (v) Fmoc-NMe-*D*-Leu (4 equiv.), HATU (4 equiv.), DIPEA (8 equiv.), RT, 2 × 1 day; (vi) 20% piperidine in DMF, RT, 2 × 30 s; (vii) HFIP/CH₂Cl₂ (1 : 4), RT, 1 h; (viii) PyBOP (3 equiv.), DIPEA (5 equiv.), DMF, RT, 1 day; (ix) 90% formic acid in water, RT, 40 min; (x) Pd-C, H₂, MeOH, RT, 90 min.



Scheme 6 Re-synthesis of **1** from the junction between *D*-Ala and NMe-*D*-Leu. Reagents and conditions: (i) Fmoc-*D*-Ala (2 equiv.), DIPEA (5 equiv.), CH₂Cl₂, RT, 1 h; (ii) iterative Fmoc-SPPS ((a) 20% piperidine in DMF, RT, 2 × 5 min; (b) Fmoc-AA-OH (4 equiv.), HATU (4 equiv.), DIPEA (8 equiv.), DMF, RT, 1 h); (iii) 20% piperidine in DMF, RT, 2 × 5 min; (iv) **3** (4 equiv.), DIC (4 equiv.), 6-Cl-HOBt (8 equiv.), DMF, RT, 24 h; (v) Fmoc-NMe-*D*-Leu (4 equiv.), HATU (4 equiv.), DIPEA (8 equiv.), RT, 2 × 1 day; (vi) HFIP/CH₂Cl₂ (1 : 4), RT, 1 h; (vii) HATU (3 equiv.), HOAt (3 equiv.), DIPEA (5 equiv.), DMF, RT, 1 day; (viii) 90% formic acid in water, RT, 40 min; (ix) Pd-C, H₂, MeOH, RT, 90 min.



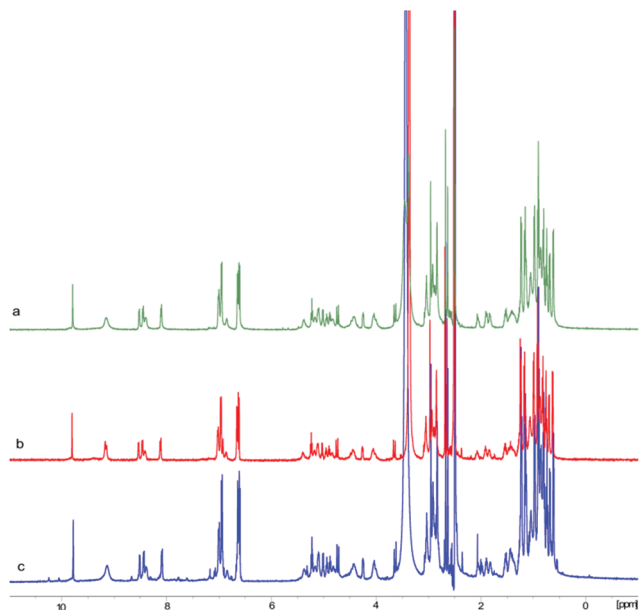


Fig. 4 Stacked ^1H -NMR (500 MHz, $\text{DMSO}-d_6$) spectra of **1** prepared from three different cyclization sites. (a) ^1H -NMR spectrum of **1** synthesized via linear peptide **4**; (b) ^1H -NMR spectrum of **1** synthesized via linear peptide **29**; (c) ^1H -NMR spectrum of **1** synthesized via linear peptide **36**. The slight differences in the three spectra are likely due to impurities from the purification steps.

Conclusions

In conclusion, the proposed structure of talarolide A (**1**) was successfully synthesized using a combination of Fmoc-SPPS and solution-phase macrolactamization. Interestingly, a novel on-resin reduction of the *N*-benzyloxy glycine residue was identified during the synthesis, which led to a low-yielding synthesis of the initial linear peptide. This side reaction was circumvented by repositioning the *N*-benzyloxy glycine residue to the middle of the linear peptide sequence. In this case, after cyclization of the linear peptide and removal of protecting groups, **1** was successfully obtained in good yield. Unfortunately, the ^1H and ^{13}C NMR spectra of the synthetic talarolide A (**1**) did not match those of the natural product and revealed the existence of two different conformers. Subsequent 2D-NMR analysis of **1**, together with the re-synthesis of **1** using two different cyclization points, fully supported **1** as the structure of our synthetic talarolide A (**1**) and also pinpointed the structural difference between the two coexisting conformers. The work reported herein suggests that further studies are required to establish the structural difference between synthetic and natural talarolide A.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors are grateful for financial support from the Maurice Wilkins Centre for Molecular Biodiscovery and a PhD scholarship from the China Scholarship Council (S. Z.).

Notes and references

- H. Drechsel and G. Jung, *J. Pept. Sci.*, 1998, **4**, 147–181.
- M. Igarashi, R. Sawa, N. Kinoshita, H. Hashizume, N. Nakagawa, Y. Homma, Y. Nishimura and Y. Akamatsu, *J. Antibiot.*, 2008, **61**, 1881–1469.
- H. Hashizume, R. Sawa, K. Yamashita, Y. Nishimura and M. Igarashi, *J. Antibiot.*, 2017, **70**, 699.
- M. A. Goetz, C. D. Schwartz, L. R. Koupal, J. M. Liesch, O. D. Hensens, R. Freidinger, P. S. Anderson, D. J. Pettibone and B. H. Woodruff, *Merck & Co Inc (US)*, EP0256847A2, 1988.
- H. Hashizume, H. Adachi, M. Igarashi, Y. Nishimura and Y. Akamatsu, *J. Antibiot.*, 2010, **63**, 279.
- K. Umezawa, K. Nakazawa, T. Uemura, Y. Ikeda, S. Kondo, H. Naganawa, N. Kinoshita, H. Hashizume, M. Hamada and T. Takeuchi, *Tetrahedron Lett.*, 1998, **39**, 1389–1392.
- D. V. Patel, M. G. Young, S. P. Robinson, L. Hunihan, B. J. Dean and E. M. Gordon, *J. Med. Chem.*, 1996, **39**, 4197–4210.
- E. Bourdel, S. Doulut, G. Jarretou, C. Labbe-Jullie, J. A. Fehrentz, O. Doumbia, P. Kitabgi and J. Martinez, *Int. J. Pept. Protein Res.*, 1996, **48**, 148–155.
- M. Marastoni, M. Bazzaro, S. Salvadori, F. Bortolotti and R. Tomatis, *Bioorg. Med. Chem.*, 2001, **9**, 939–945.
- A. Bianco, C. Zabel, P. Walden and G. Jung, *J. Pept. Sci.*, 1998, **4**, 471–478.
- T. Kolasa, *Tetrahedron*, 1983, **39**, 1753–1754.
- A. Bianco, D. Kaiser and G. Jung, *Chem. Biol. Drug Des.*, 1999, **54**, 544–548.
- P. Dewapriya, P. Prasad, R. Damodar, A. A. Salim and R. J. Capon, *Org. Lett.*, 2017, **19**, 2046–2049.
- S. B. Singh, J. Odingo, M. A. Bailey, B. Sunde, A. Korkegian, T. O'Malley, Y. Ovechkina, T. R. Ioerger, J. C. Sacchettini and K. Young, *bioRxiv, Microbiol.*, 2018, 279307.
- Y. M. Elbatrawi, C. W. Kang and J. R. Del Valle, *Org. Lett.*, 2018, **20**, 2707–2710.
- C. J. White and A. K. Yudin, *Nat. Chem.*, 2011, **3**, 509.
- A. K. Ghosh and C.-X. Xu, *Org. Lett.*, 2009, **11**, 1963–1966.
- S. Zhang, L. M. De Leon Rodriguez, E. Lacey, A. M. Piggott, I. K. Leung and M. A. Brimble, *Eur. J. Org. Chem.*, 2017, 149–158.
- M. Teixidó, F. Albericio and E. Giralt, *J. Pept. Res.*, 2005, **65**, 153–166.
- X. Hu, J. Zhu, S. Srivathsan and D. Pei, *Bioorg. Med. Chem. Lett.*, 2004, **14**, 77–79.
- Y. Ye, M. Liu, J. L. K. Kao and G. R. Marshall, *Pept. Sci.*, 2003, **71**, 489–515.



- 22 H. C. J. Ottenheijm and J. D. M. Herscheid, *Chem. Rev.*, 1986, **86**, 697–707.
- 23 A. Siow, G. Opiyo, I. Kavianinia, F. F. Li, D. P. Furkert, P. W. R. Harris and M. A. Brimble, *Org. Lett.*, 2018, **20**, 788–791.
- 24 R. E. Steiger, *J. Biol. Chem.*, 1944, **153**, 691–692.
- 25 H. C. J. Ottenheijm, R. Plate, J. H. Noordik and J. D. M. Herscheid, *J. Org. Chem.*, 1982, **47**, 2147–2154.
- 26 T. Kolasa, *Synthesis*, 1983, 539–539.
- 27 J. D. M. Herscheid, R. J. F. Nivard, M. W. Tijhuis, H. P. H. Scholten and H. C. J. Ottenheijm, *J. Org. Chem.*, 1980, **45**, 1880–1885.
- 28 H. Kaur, A. M. Heapy, R. Kowalczyk, Z. Amso, M. Watson, J. Cornish and M. A. Brimble, *Tetrahedron*, 2014, **70**, 7788–7794.
- 29 S. Zhang, Z. Amso, L. M. De Leon Rodriguez, H. Kaur and M. A. Brimble, *J. Nat. Prod.*, 2016, **79**, 1769–1774.
- 30 Y. Takeuchi and G. R. Marshall, *J. Am. Chem. Soc.*, 1998, **120**, 5363–5372.
- 31 V. Dupont, A. Lecoq, J. P. Mangeot, A. Aubry, G. Boussard and M. Marraud, *J. Am. Chem. Soc.*, 1993, **115**, 8898–8906.
- 32 M. Akiyama, K. Lesaki, A. Katoh and K. Shimizu, *J. Chem. Soc., Perkin Trans. 1*, 1986, 851–855.

