Bicyclic Pyrrolidine-Isoxazoline γ Amino Acid: A Constrained Scaffold for Stabilizing α-Turn Conformation in Isolated Peptides

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Unnatural amino acids have tremendously expanded the folding possibilities of peptides and peptide mimics. While α,α-disubstituted and β-amino acids are widely studied, γ-derivatives have been less exploited. Here we report the conformational study on the bicyclic unnatural γ amino acid, 4,5,6,6a-tetrahydro-3aH-pyrrolo[3,4-d]isoxazole-3-carboxylic acid 1. In model peptides, the (+)-(3aR,6aS)-enantiomer is able to stabilize α-turn conformation when associated to glycine, as showed by 1H-NMR, FT-IR, and circular dichroism experiments, and molecular modeling studies. α-turn is a structural motif occurring in many biologically active protein sites, although its stabilization on isolated peptides is quite uncommon. Our results make the unnatural γ-amino acid 1 of particular interest for the development of bioactive peptidomimetics.

Keywords: unnatural γ-amino acids, peptidomimetic, isoxazoline, α-turn, metadynamic studies, conformational analysis

INTRODUCTION

Amino acids are the key building blocks of proteins and biomolecules and are widely exploited in different applications, from pharmaceutical chemistry and biomedicine to material science, optoelectronics and catalysis (Zhang et al., 2012; Mikhalevich et al., 2017; Solomon et al., 2017; López-Andarias et al., 2018; Raymond and Nilsson, 2018). The insertion of unnatural amino acids (UAAs) in peptides and peptide mimics could add specific features to these molecules, such as proteolytic stability, active functional groups and new reactivity (Clerici et al., 2016; Pellegrino et al., 2016, 2017; Bucci et al., 2017). α,α disubstituted and β- homologs of natural amino acids have been particularly studied during the years for their ability to introduce conformational constrains in peptides and to stabilize specific secondary structures (Bonetti et al., 2015; Pellegrino et al., 2015; Fanelli et al., 2017; Kobayashi et al., 2017). On the other hand, γ UAAs provide a further opportunity to engineer the available backbone through the incorporation of an additional methylene group...
(Vasudev et al., 2011; Sohora et al., 2018). Recent studies report that γ-peptides are able to form helices, sheets and turns, whose conformational stability is increased by introducing substituents on the backbone chain (Bouillère et al., 2011). Moreover, cyclic γ UAAls, as gabapentin (Gpn) (Chatterjee et al., 2009; Konda et al., 2018) and γ-cyclohexyl amino acid (Guo et al., 2012) are able to stabilize both turn and helix conformation in oligomers and in mixed α-γ and β-γ sequences. On the other hand, the use of bicyclic γ amino acids for the stabilization of the peptide structure is much less investigated (Machetti et al., 2000).

In this work, we investigated the conformational behavior of both the enantiomers of the bicyclic unnatural γ amino acid 4,5,6,6a-tetrahydro-3aH-pyrrolol[3,4-d]isoxazole-3-carboxylic acid 1, obtained starting from the corresponding ethyl esters recently described by us (Tamborini et al., 2015). Compound 1 is a conformationally constrained dipeptide analog and, in principle, it could substitute two amino acids in a peptide chain. The presence of the constrained bicyclic system could induce specific secondary structure allowing a proper orientation of the peptide arms at C- and N- termini. Furthermore, the presence of the isoxazoline ring, a core often found in biologically active compounds, could be particularly useful for future applications in the pharmaceutical field. In fact, isoxazoline derivatives are important scaffolds found in many naturally occurring and biologically active compounds possessing a wide range of bioactivities, such as antibacterial, antifungal, antiparasitic (Conti et al., 2011; Bruno et al., 2014; Pinto et al., 2016a), anticancer (Castellano et al., 2011; Kaur et al., 2014), anti-inflammatory and anticonvulsant activity (Sperry and Wright, 2005; Pinto et al., 2011, 2016b). Isoxazolines are also considered to be important precursors for the synthesis of β-hydroxyketones (Kozikowski and Park, 1990; Tsantali et al., 2007), β-aminoalcohols (Fuller et al., 2005), isoxazolidines (Itoh et al., 2002), and many other valuable compounds. Recently, peptidomimetics containing the isoxazoline ring have been reported as β-turn mimics (Bucci et al., 2018; Memeo et al., 2018).

Starting from the (−)-(3aS,6aR)-1 and (+)-(3aR,6aS)-1 enantiomers, model peptides containing Leu-Val dipeptide at C-terminus and variable sequences at N-terminus were prepared. Their conformational behavior was investigated by NMR spectroscopy, FT-IR, circular dichroism, and molecular modeling. Our results indicated that (+)-(3aR,6aS)-1 enantiomer, in combination with glycine, is effective in stabilizing the α-turn conformation in peptides (Figure 1). This structural motif occurs quite often in many key sites of proteins, such as enzyme active site, and metal binding domains (Wintjens et al., 1996), although few molecules are known to mimic or stabilize it on isolated peptides (Kelso et al., 2004; Hoang et al., 2011; Krishna et al., 2014; Wang et al., 2018). Our results make thus the unnatural γ-amino acid 1 of particular interest for future development of bioactive peptidomimetics.

**Abbreviations:** ACN, Acetonitrile; Boc, tert-butyloxycarbonyl; DIEA, N,N-Disopropylethylamine; DMF, Dimethylformamide; EDC, 1-Ethyl-3-(3-dimethylaminopropyl)carbodiimide; Gly, Glycine; HOBt, Hydroxybenzotriazole; Leu, Leucine; Phe, Phenylalanine; TFA, Trifluoroacetic acid; Val, Valine.

### MATERIALS AND METHODS

#### Materials

Chemicals were obtained from Zentek (Italy) and used without further purification. Melting points were determined in a Stuart Scientific melting point apparatus in open capillary tubes and are certified. ESI mass spectra were recorded on an LCQESI MS was recorded on a LCQ Advantage spectrometer from Thermo Finnigan and a LCQ Fleet spectrometer from Thermo Scientific. CD experiments were carried out on a Jasco J-810 instrument. Spectra were obtained from 195 to 250 nm with a 0.1 nm step and 1 s collection time per step, taking three averages. The CD spectra were plotted as mean residue ellipticity \( \theta \) (degree \( \times \) cm\(^2\) \times dmol\(^{-1}\)) vs. wave length \( \lambda \) (nm). Noise-reduction was obtained using a Fourier-transform filter program from Jasco. The NMR spectroscopic experiments were carried out on a Varian OXFORD 300 MHz (300 and 75 MHz for \( ^1\)H and \( ^{13}\)C, respectively). Chemical shifts, \( \delta \), are given in ppm relative to the CHCl\(_3\) internal standard, and the coupling constants, \( J \), are reported in hertz (Hz).

#### Methods

**Synthesis of (−)-(3aS,6aR)-1 and (+)-(3aR,6aS)-1**

Compound (−)-2 or (+)-2 (200 mg, 0.70 mmol) was dissolved in MeOH (3.0 mL) and treated with 1N NaOH aqueous solution (1.4 mL). The mixture was stirred at room temperature for 1 h and the disappearance of the starting material was monitored by TLC (Cyclohexane/EtOAc 7:3). The mixture was diluted in water (20 mL), made acidic with 2N aqueous HCl and extracted with EtOAc. The organic phase was dried over Na\(_2\)SO\(_4\) and after evaporation of the solvent, the acid derivate (−)-(3aS,6aR)-1 or (+)-(3aR,6aS)-1 (170 mg, 95%) was obtained as a white solid.

(−)-(3aS,6aR)-1: \( R_f \left(\text{CH}_2\text{Cl}_2/\text{MeOH} 9:1 + 1\% \text{AcOH}\right): 0.40; [\alpha]_D^{20} = -192.2 \text{ (c: 0.55 in MeOH)} \); mp = dec. \( T > 137^\circ\text{C} \); \( ^1\)H NMR (300 MHz, CD\(_3\)OD): \( \delta 5.35 \) (dd, \( j = 4.8, 9.2, 1H \)); 4.10 (ddd, \( j = 2.2, 9.2, 9.2, 1H \)); 3.86–3.96 (m, 2H); 3.38–3.50 (m, 2H); 3.12–3.25 (m, 2H); 2.97–3.01 (m, 1H); 2.82–2.97 (m, 2H); 2.55–2.66 (m, 1H); 1.44 (s, 9H) ppm; \( ^{13}\)C NMR (75 MHz, CD\(_3\)OD): \( \delta 161.28, 154.77, 152.95, 87.62, 80.33, 53.06, 50.93, 49.10, 27.12 (3C) \) ppm.

**HRMS (ESI) [M + Na]\(^+\) calculated for C\(_{11}\)H\(_{16}\)N\(_2\)O\(_2\)Na: 279.0957, found: 279.0950.**

(+)-(3aR,6aS)-1: [\( \alpha \)]\(_D^{20}\) = +192.5 (c: 0.57 in MeOH).

**General Procedure for Coupling Reaction**

Free carboxylic acid (1 equiv.) was dissolved in CH\(_2\)Cl\(_2\) (0.01M solution) and the mixture was cooled to 0°C. HOBt (1.1 equiv.) and EDC (1.1 equiv.) were added. After 1 h, free amino compound (1 equiv.) was added, followed by the addition of DIEA (2 equiv.). The reaction mixture was stirred at room temperature for 24 overnight (TLC analysis). The organic layer was washed with a 5% solution of KHSO\(_4\) (3 times), with a saturated solution of NaHCO\(_3\) (3 times) and with brine (1 time). The organic phase was dried over Na\(_2\)SO\(_4\) and the solvent was removed under vacuum. The products were purified by column flash chromatography on silica gel (hexane/ethyl acetate gradient) followed by a crystallization from a mixture of ethyl acetate/hexane.
Steinke and Kula (1990)

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γ

N-Boc-(–)-490.2642, found: 490.2635.

28.31, 24.82, 22.91, 21.92, 19.27, 17.93 ppm.

153.93, 87.40, 80.43, 58.24, 53.26, 52.30, 50.53, 49.32, 40.87, 30.53, 1.49 (s, 9H), 0.99 – 0.92 (m, 12H) ppm.

13C NMR (75 MHz, CDCl3): δ 173.04, 171.32, 159.44, 154.28, 153.93, 87.40, 80.43, 58.24, 53.26, 52.30, 50.53, 49.32, 40.87, 30.53, 28.31, 24.82, 22.91, 21.92, 19.27, 17.93 ppm.

HRMS (ESI) [M + Na]+ calculated for C$_{31}$H$_{46}$N$_6$O$_7$Na: 637.3326, found: 637.3332.

**N-Boc-(-)-Δ$^2$-isox-Leu-Val-CONH$_2$ (4)**

1H NMR (300 MHz, CDCl$_3$): δ 7.11 (bs, 1H), 6.88 (d, J = 8.8 Hz, 1H), 6.19 (bs, 1H), 5.72 (bs, 1H), 5.34–5.29 (m, 1H), 4.56–4.48 (m, 1H), 4.31 (dd, J = 8.2; 6.9 Hz, 1H), 4.17–3.83 (m, 3H), 3.56–3.42 (m, 2H), 2.23–2.07 (m, 1H), 1.85–1.61 (m, 3H), 1.43 (s, 9H), 0.99 – 0.92 (m, 12H) ppm.

13C NMR (75 MHz, CDCl$_3$): δ 173.04, 171.32, 159.44, 154.28, 153.93, 87.40, 80.43, 58.24, 53.26, 52.30, 50.53, 49.32, 40.87, 30.53, 28.31, 24.82, 22.91, 21.92, 19.27, 17.93 ppm.

HRMS (ESI) [M + Na]+ calculated for C$_{31}$H$_{46}$N$_6$O$_7$Na: 637.3326, found: 637.3332.

**N-Boc-(+)-Δ$^2$-isox-Leu-Val-CONH$_2$ (5)**

1H NMR (300 MHz, CDCl$_3$): δ 7.05 (bs, 1H), 6.73 (d, J = 7.9 Hz, 1H), 6.13 (bs, 1H), 5.69 (bs, 1H), 5.37–5.28 (m, 1H), 4.56–4.47 (m, 1H), 4.36–4.27 (m, 1H), 4.18–4.07 (m, 1H), 3.99–3.77 (m, 2H), 3.61–3.41 (m, 2H), 2.28–2.10 (m, 1H), 1.78–1.57 (m, 3H), 1.49 (s, 9H), 1.01 – 0.90 (m, 12H) ppm.

13C NMR (75 MHz, CDCl$_3$): δ 173.16, 171.54, 159.33, 154.16, 153.96, 80.34, 60.38, 58.16, 53.27, 52.25, 49.41, 41.02, 30.67, 28.32, 24.83, 22.87, 21.99, 19.20, 18.02 ppm.

HRMS (ESI) [M + Na]+ calculated for C$_{22}$H$_{37}$N$_5$O$_6$Na: 490.2642, found: 490.2635.

**NH-Boc-Phe-(+)-Δ$^2$-isox-Leu-Val-CONH$_2$ (9) – Mixture of two conformers**

1H NMR (300 MHz, CD$_2$CN): δ 7.33–7.05 (m, 6H), 7.04–6.91 (m, 1H), 6.70 (d, J = 8.5 Hz, 1H), 6.13–5.91 (m, 2H), 5.46 (d, J = 8.5 Hz, 1H), 5.36–5.10 (m, 2H), 4.87–4.71 (m, 1H), 4.67–4.43 (m, 1H), 4.38–4.23 (m, 2H), 4.23–3.69 (m, 4H), 3.69–3.41 (m, 1H), 3.21 (dd, J = 14.3; 5.0 Hz, 1H), 3.10–2.79 (m, 2H), 2.54 (dd, J = 14.3; J = 5.0 Hz, 1H), 2.30–1.99 (m, 1H), 1.89–1.55 (m, 3H), 1.49 (m, 9H), 1.31 (m, 12H) ppm.

13C NMR (75 MHz, CD$_2$CN): δ 173.14, 172.90, 171.92, 171.23, 170.66, 170.49, 159.68, 158.86, 155.92, 155.18, 153.77, 153.59, 136.00, 129.63, 129.43, 129.15, 128.60, 128.41, 127.14, 127.05, 86.96, 85.53, 79.99, 79.88, 58.23, 58.73, 54.13, 53.47, 52.97, 52.19, 51.05, 48.82, 41.04, 40.85, 39.32, 30.88, 30.66, 28.45, 28.30, 25.05, 24.81, 22.91, 22.01, 21.51, 19.28, 17.98, 17.84 ppm.

HRMS (ESI) [M + Na]+ calculated for C$_{31}$H$_{46}$N$_6$O$_7$Na: 637.3326, found: 637.3332.

**NH-Boc-Phe-Gly-(+)-Δ$^2$-isox-Leu-Val-CONH$_2$ (10)**

1H NMR (300 MHz, CD$_2$CN): δ 7.47 (d, J = 6.9 Hz, 1H), 7.30–7.26 (m, 6H), 6.95 (d, J = 8.5 Hz, 1H), 6.72 (bs, 1H), 6.42 (bs, 1H), 5.63 (bs, 1H), 5.47–5.31 (m, 1H), 4.52–4.47 (m, 1H), 4.40–3.39 (m, 8H), 3.23–3.07 (m, 1H), 2.70–2.93 (m, 1H), 2.14–2.08 (m, 1H), 1.78–1.55 (m, 3H), 1.35 (s, 9H), 1.02–0.77 (m, 12H) ppm.

13C NMR (75 MHz, CD$_2$CN): δ 173.43, 171.63, 167.09, 159.52, 154.34, 137.64, 129.27, 128.31, 126.53, 109.99, 87.41, 86.02, 79.09,
General Procedure for N-Boc-Deprotection
N-Boc protected compound (1 equiv.) was dissolved in CH₂Cl₂ (0.01M) and the solution cooled at 0°C. TFA (50% v/v) was added dropwise, the solution was warmed up at room temperature and was stirred for 2 h. The solvent was removed under vacuum with the obtainment of the trifluoroacetate salt that was directly used in the next coupling step.

RESULTS AND DISCUSSION

The two enantiomers of compound 1 were obtained starting from the corresponding ethyl esters recently described by us (Tamborini et al., 2015). Racemate (±)-2 (Conti et al., 2006) was synthesized in a flow chemistry reactor exploiting the 1,3-dipolar cycloaddition reaction of N-Boc-3-pyrrole with ethoxybenzyl formonitrile oxide. An excellent enantiomeric separation (e.e. >99%) of the racemate was achieved by semi-preparative chiral HPLC. Alkaline hydrolysis of (+)-(3aS6aR)-2 and (+)-(3aR6aS)-2 gave the desired (−)-(3aS6aR)-1 and (+)-(3aR6aS)-1, respectively (Scheme 1).

The ability of compound 1 to stabilize secondary structures was evaluated in model peptides containing (L)-Phe at N-terminus and (L)-Leu-(L)-Val sequence at C-terminus and (Schemes 2, 3). This last dipeptide was chosen as it normally adopts extended conformation in solution. Compound 1 was used in both the 3aS6aR and 3aR6aS

| Product | Yield (%) |
|---------|-----------|
| 3a      | 95        |
| 6       | 95        |
| 7       | 95        |

\(^a\) see Smith and Spackman (1955).
stereochemistries [(–)-1 and (+)-1, respectively], as a different effect on peptide conformations could be expected depending on the stereochemistry of the unnatural amino acids (Pellegrino et al., 2012).

Firstly, Leu-Val dipeptide 3 was prepared starting from (1)-Valinamide and NH-Boc-(1)-Leu, according to the general coupling procedure, followed by N-terminus deprotection in TFA (91% overall yield). Diastereoisomeric dipeptides 4 and 5 were then achieved in good yields (65% and 63%, respectively) by a coupling reaction of (–)-1 or (+)-1 with dipeptide 3 (Scheme 2).

Both 4 and 5 compounds were completely characterized by NMR spectroscopy (CDCl₃, see SI for complete proton assignment). In both tripeptides, no significant Noesy effects between Leu-Val dipeptide and the scaffold were detected. Furthermore, in variable temperature experiments, the obtained $\Delta \delta/\Delta T$ is higher than 5 ppb/K for all the amide protons, indicating the absence of H-bonds stabilizing a particular conformation. The $J_{NH-\text{CH}_\text{a}}$ value for NH-Val is of 8.8 Hz and 7.9 Hz, for 4 and 5, respectively (NH-Leu appears as a broad signal for each compound). From these findings, we hypothesized that both 4 and 5 adopt an extended conformation and that the isoxazoline scaffold is not interacting with Leu-Val dipeptide.

The peptide chain was then elongated at N-terminus, through Boc-deprotection and coupling with (1)-Phe. Diastereoisomeric dipeptides 8 and 9 were achieved in good overall yields (73% and 76%, respectively) (Scheme 3). From NMR studies, it was found that both model peptides 8 and 9 are present as a mixture of conformers in 2:1 ratio in CDCl₃ solution. The presence of these two conformers is probably due to the low-energy barrier $cis/trans$ isomerization of the tertiary amide on the pyrrolidine as it is frequently observed on the acylated proline and in the case of tertiary cyclic amides (Laursen et al., 2013; Pellegrino et al., 2014). Regarding compound 8, this rotation led to a splitting of the (1)-Phe proton signals only. Furthermore, no significative Noesy effects were detected, and, in variable temperature experiments, the obtained $\Delta \delta/\Delta T$ coefficient is high for all the amide protons. Taking all these data together, we can assume that Phe and Leu-Val dipeptide are oriented in opposite directions and that the 3aS6aR stereochemistry of the scaffold is not effective in inducing specific secondary structures when used in combination with (1)-α-amino acids.

A different scenario was observed for compound 9. Its two rotamers are indeed characterized by different chemical shifts, suggesting that the isomerization of the tertiary amide leads to two different structures conformations influencing the entire molecule. Unfortunately, due to several overlapping signals, it was not possible to assessing significative Noesy proximities. Variable temperature experiments were thus done, in order to investigate if the NH were involved in hydrogen bonds and, as a consequence, if the two isomers of 9 were characterized by a stable conformation in solution. In the case of the major isomer, the obtained $\Delta \delta/\Delta T$ is around 4 ppb/K for NH-Phe, indicating that this proton could be involved in a weak/medium hydrogen bond. All the other amide protons had higher $\Delta \delta/\Delta T$. On the
other hand, in the minor conformer the obtained Δh/ΔT is of around 2 ppb/K for NH-Val, indicating that this proton is involved in a strong hydrogen bond. Metadynamic studies were then performed on the cis/trans tertiary amide conformers of compound 9 (Figure 2).

In particular, we performed two 50 ns long metadynamics simulations, using the distance between Boc quaternary carbon and Cβ Val as collective variable (CV). This geometric parameter was selected as a suitable CV because it discriminates well between closed and extended states of the peptide. In this way, it could be possible to evaluate if the two conformers had an intrinsic tendency toward turn conformation. The cis isomer showed a broad free energy minimum corresponding to CV values between 5 and 8 Å, while the trans isomer showed much higher free energy values in this region, exhibiting a shallow minimum around 10–15 Å (Figure 3).

This different behavior could be indicative of a preference of cis-9 toward a more closed conformation, although from these data it was not possible to make any further assumptions. For this reason, we envisaged that the introduction of a Gly residue between Phe and isoxazoline scaffold could increase the conformational flexibility of the N-terminus peptide arm favoring its interaction with the C-terminus part of the molecule. Compound 7 was thus elongated at N-terminus, through coupling with N-Boc-(L)-Phe-Gly-OH dipeptide using general coupling reaction conditions to obtain peptide 10.
(65%, Scheme 3). The NMR study on 10 (CD$_3$CN, see SI for complete assignment), showed that it is present in solution as a single stable conformer (only trace amounts of a second isomer were detected). A complete set of NH$_i$-NH$_{i+1}$ Noesey proximities, whose calculated distances ranged from 2.68 to 2.96 Å, were observed (Figure 4 on the top right). Furthermore, long range Noesy effects involving H$_\alpha$Gly and NHVal (3.03 Å) and H$_\beta$Leu (2.73 Å) were found (Figure 4 on the bottom).

In variable temperature experiments (Figure 5), the obtained $\Delta\delta/\Delta T$ is of around 2 ppb/K for NH-Val, while NH-Leu and one of the C-terminus amide protons possess $\Delta\delta/\Delta T$ of around 3 and 4 ppb/K, respectively. The presence of hydrogen bonds was also confirmed by FT-IR analysis. N-H stretching bands A and B (3,500–3,000 cm$^{-1}$ region, Figure S20) are indeed downshifted as frequently observed in intramolecular hydrogen bonded conformations (Tonan and Ikawa, 1996; Barth, 2007). Furthermore, the deconvolution of the amide I band (1,700–1,600 cm$^{-1}$) showed the presence of a band at 1,655 cm$^{-1}$ (Figure S20), typical of $\alpha$ structures (Kong and Yu, 2007).

In order to gain more information on the conformation of compound 10, far-UV circular dichroism (CD) analysis in CH$_3$CN (0.1 mM solution) was then performed. In Figure 6 the obtained CD spectrum is reported.

It showed a maximum at around 250 nm due to the strong absorption of the isoxazoline ring, as observed by Memeo et al. (2018) on similar compounds (see also the UV spectrum in the SI). In the amide bond region, two negative minima at 225 and 208 nm, with the same intensity, and a slightly positive band at 200 nm were also observed. The presence of the exciton splitting of the $\pi \rightarrow \pi^*$ transition band suggested that a $\alpha$-turn conformation was present (Wang et al., 2018), although in short
In order to completely elucidate the conformation of peptide 10, a restrained molecular dynamic was finally performed. The restraints were based on the obtained NOESY values. When the hydrogen was not uniquely defined, like in H_αGly and H_βLeu, we used the closest carbon, e.g., C_αGly and C_βLeu to implement distance restraints (Figure S3). The analysis showed that C_αGly-C_βLeu (A) and C_αGly-H_NHVal (B) are mutually exclusive in the trans isomer but not in the cis one as shown in Figure S1. Furthermore, H-bond analysis resulted in the observation that a H-bond is between CO-Gly and NH-Val. In Figure 7 the most representative structure is reported.

Taking together both experimental and molecular modeling data, we can assume that, in solution, compound 10 effectively adopts a α-turn conformation. This motif is normally formed by 5 α-amino acids and is characterized by a H-bond between i and i+4 residues. In our case, the isoxazoline scaffold replaces two of the three core amino acids (Figure 1) and the H-bond is formed by the CO-Gly and NH-Val, leading to the formation of a 12 member pseudo-cycle. Furthermore, the overall structure of the peptidomimetic is reinforced by medium-weak H-bonds involving NH-Leu and one C-terminus-NH_2 as evicted from NMR data. The presence of a α-turn conformation is also confirmed by the computational analysis of the dihedral angles of the residue i+3 and of the distances between residues i and i+4 (Table S9), as their values rely within the α-turn parameters (Pavone et al., 1996).

In conclusion, as α-turns are often found on biologically active sites and few molecules have been reported to mimic or stabilize them, the ability of compound (+)-1 to stabilize α-turn conformation on isolated peptide is particularly important in view of future biological applications.

**DATA AVAILABILITY**

All datasets generated for this study are included in the manuscript and/or the supplementary files.

**AUTHOR CONTRIBUTIONS**

FO and RB equally contributed to this work. The project was conceived by SaP, LT, and AP. SaP designed and coordinated the research. LT and RB chemically synthesized and analyzed the materials, RB performed NMR, FT-IR and CD experiments, FO and StP performed molecular modeling. SaP, StP, LT, and AP analyzed and compiled the data and co-wrote the manuscript. The final manuscript was read and approved by all the authors.
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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fchem.2019.00133/full#supplementary-material
Mikhalevich, V., Craciun, I., Kyropoulou, M., Palivan, C. G., and Meier, W. (2017). Amphiphilic peptide self-assembly: expansion to hybrid materials. Biomacromolecules 18, 3471–3480. doi: 10.1021/acs.biomac.7b00764

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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