


 Cite this: *RSC Adv.*, 2022, 12, 4640

# Controlling the conformational stability of coiled-coil peptides with a single stereogenic center of a peripheral $\beta$ -amino acid residue†

 Monika Szefczyk, \*<sup>a</sup> Katarzyna Ożga, <sup>a</sup> Magda Drewniak-Świtalska,<sup>a</sup>  
 Ewa Rudzińska-Szostak, <sup>a</sup> Rafał Hołubowicz,<sup>b</sup> Andrzej Ozyhar <sup>b</sup>  
 and Łukasz Berlicki \*<sup>a</sup>

The key issue in the research on foldamers remains the understanding of the relationship between the monomers structure and conformational properties at the oligomer level. In peptidomimetic foldamers, the main goal of which is to mimic the structure of proteins, a main challenge is still better understanding of the folding of peptides and the factors that influence their conformational stability. We probed the impact of the modification of the peptide periphery with *trans*- and *cis*-2-aminocyclopentanecarboxylic acid (ACPC) on the structure and stability of the model coiled-coil using circular dichroism (CD), analytical ultracentrifugation (AUC) and two-dimensional nuclear magnetic resonance spectroscopy (2D NMR). Although, *trans*-ACPC and *cis*-ACPC-containing mutants differ by only one peripheral stereogenic center, their conformational stability is strikingly different.

 Received 7th January 2022  
 Accepted 31st January 2022

DOI: 10.1039/d2ra00111j

[rsc.li/rsc-advances](http://rsc.li/rsc-advances)

## Introduction

The main inspiration for the development of foldamers, artificial oligomers with a strong tendency to adopt a well-defined thermodynamically stable conformation in solution, were proteins. It was assumed that it might be possible to design similar but unnatural oligomers that fold into compact and specific conformations leading to new types of molecules with interesting properties and useful applications.<sup>1</sup> Much research has been undertaken on artificial scaffolds capable of mimicking naturally occurring structural motifs, *i.e.* coiled-coils.<sup>2,3</sup> The “bottom-up” approaches use *de novo* designed foldameric helices to obtain higher-order structures, usually by the segregation of hydrophilic and lipophilic side chains in the helical conformation.<sup>4</sup> In the “top-down” methods, a previously characterized  $\alpha$ -amino acid sequence of known secondary/tertiary/quaternary structure is applied directly in the design of peptides with nonnatural backbones intended to form helix bundles.<sup>5–7</sup> The crucial step in the design is the identification of new backbones with well-defined structural preferences. Oligo( $\beta$ -amino acids), named  $\beta$ -peptides, were first proved to meet

this criterion and have been widely studied *i.e.* in terms of folding properties.<sup>8–10</sup> In addition,  $\beta$ -peptides show improved stability in enzymatic degradation and have various biological activities, *e.g.* antimicrobial.<sup>11</sup> It was soon proved that a very useful approach for the development of foldamers is the introduction of unnatural amino acids into  $\alpha$ -peptides in the process called foldamerization.<sup>12,13</sup> In terms of peptidomimetics, apart from widely studied foldamers based on higher homologues of  $\alpha$ -amino acids,<sup>14</sup> there have been also studied the sequences containing  $\alpha$ -aminoisobutyric acid (Aib),<sup>15–17</sup> oligooureas,<sup>18</sup> azapeptides,<sup>19</sup>  $\alpha$ -hydrazido-peptides,<sup>20</sup> polyamides,<sup>21</sup> and others.<sup>22–24</sup> Foldamerization of sequences using  $\beta$ -amino acids shows two main advantages, namely, it improves both the conformational and proteolytic stability of peptides.<sup>25–27</sup> Importantly, retaining activity<sup>28</sup> and binding affinity<sup>29–31</sup> of the modified peptides after  $\alpha \rightarrow \beta$  replacement is usually possible. A wide range of possible applications of the reported  $\alpha$ , $\beta$ -peptides prompted extensive studies on the influence of constrained residues on secondary structure formation, mainly in terms of improving their conformational stability and chosen properties.<sup>32–34</sup>

The influence of constrained  $\beta$ -amino acid residues on secondary structure formation could also potentially have a destructive effect. Analysis of the analogues with conformational stability significantly lower than that of the original oligomer could provide knowledge of the folding propensities of the studied oligomers as well as indicate properties of individual monomers. It is worth noting that the reason for the conformational instability upon substitution could be at least of two types: (a) steric impairment of the side chain with the rest of

<sup>a</sup>Department of Bioorganic Chemistry, Faculty of Chemistry, Wrocław University of Science and Technology, Wybrzeże Wyspiańskiego 27, 50-370 Wrocław, Poland. E-mail: monika.szefczyk@pwr.edu.pl; lukasz.berlicki@pwr.edu.pl

<sup>b</sup>Department of Biochemistry, Molecular Biology and Biotechnology, Faculty of Chemistry, Wrocław University of Science and Technology, Wybrzeże Wyspiańskiego 27, 50-370 Wrocław, Poland

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d2ra00111j





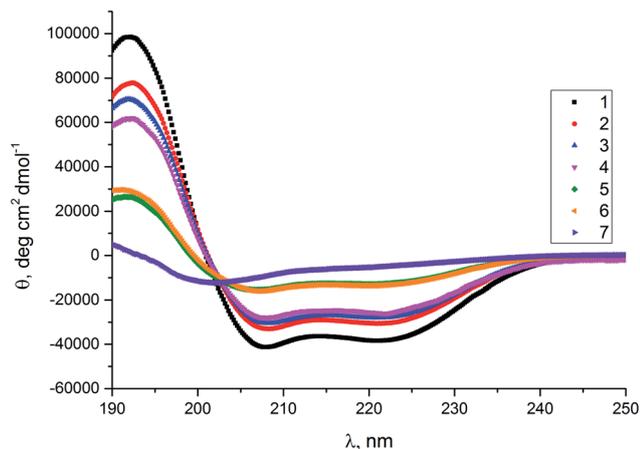


Fig. 2 CD spectra of the studied peptides in phosphate buffer.  $C_{\text{pep}} = 80 \mu\text{M}$ ,  $C_{\text{buffer}} = 0.05 \text{ M}$ ,  $\text{pH} = 7$ ,  $T = 20 \text{ }^\circ\text{C}$ .

$R(\theta_{222}/\theta_{208})$  close to 1, which confirms the presence of  $\alpha$ -helix structure.<sup>49</sup> Introduction of only one *trans*- $\beta$  residue (peptides 2 and 3) does not change the position of the minima compared to the model peptide 1 and does not have a significant influence on the  $R(\theta_{222}/\theta_{208})$  and molar ellipticity values. The  $\alpha \rightarrow \text{trans}$ -ACPC residue replacement at both ends of the sequence (peptide 4) resulted in a slight decrease in the molar ellipticity value compared to the peptides mentioned above and the position of two minima was also preserved. In the spectra of peptides 5 and 6, where a single  $\alpha \rightarrow \text{cis}$ -ACPC residue replacement was introduced, a more significant decrease in the molar ellipticity values can be observed, compared to peptide 1, and two shallow minima at  $\lambda = 208$  and  $222 \text{ nm}$ . The introduction of *cis*- $\beta$  residue at both ends (peptide 7) caused total disruption of the helix structure, since only a minimum at  $204 \text{ nm}$  is present in the CD spectrum, indicating a random coil formation.<sup>50</sup>

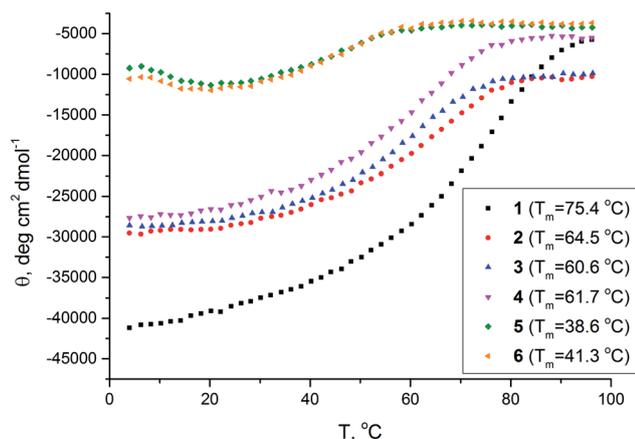


Fig. 3 Thermal unfolding curves of the studied peptides followed by CD signal at  $222 \text{ nm}$  in phosphate buffer with calculated melting temperatures indicated in the legend.  $C_{\text{pep}} = 80 \mu\text{M}$ ,  $C_{\text{buffer}} = 0.05 \text{ M}$ ,  $\text{pH} = 7$ .

Conformational stability was determined based on CD measurements in phosphate buffer and was followed in the range of  $190\text{--}240 \text{ nm}$  by a gradual increase in temperature from  $4$  to  $98 \text{ }^\circ\text{C}$  (Fig. S2†). Thermal unfolding curves followed at  $222 \text{ nm}$  were used to determine the melting temperatures (Fig. 3). The model coiled-coil peptide 1 is very stable in the phosphate buffer with a cooperative melting transition at temperature  $75 \text{ }^\circ\text{C}$ . A singular  $\alpha \rightarrow \text{trans}$ - $\beta$  residue replacement was shown to have an influence on  $\alpha, \beta$ -peptides stability, since the melting temperatures calculated for peptides 2 and 3 are lower compared to the model peptide 1, however still very high (more than  $60 \text{ }^\circ\text{C}$ ). Furthermore, the introduction of two *trans*- $\beta$  residues (peptide 4) still preserves a stable helical structure. On the other hand, even a singular peripheral  $\alpha \rightarrow \text{cis}$ - $\beta$  modification has a dramatic influence on the stability of the peptides. The melting temperatures obtained for peptides 5 and 6 are significantly lower compared to other peptides studied. Data for peptide 7 were not obtained as the CD spectrum did not indicate  $\alpha$ -helical fold.

In summary,  $\alpha \rightarrow \text{trans}$ - $\beta$  residue replacement does not significantly influence the stability of the peptides in buffer, although  $\alpha \rightarrow \text{cis}$ - $\beta$  residue replacement caused the disruption of the structure.

The peptides modified with *trans*-ACPC or *cis*-ACPC were subjected to sedimentation velocity (SV AUC) and sedimentation equilibrium (SE AUC) analytical ultracentrifugation to determine their oligomerization states and the effect of unnatural amino acids incorporation on the stability of the oligomeric forms. Peptides 1, 2, 3, and 4 existed in solution as trimers with sedimentation coefficients  $0.90\text{--}0.93 \text{ S}$ , as determined by the SV AUC (Table 1, Fig. S3†). In addition to trimers, the sedimentation coefficient distributions ( $c(s)$ ) indicate that low molecular weight components ( $0.25\text{--}0.30 \text{ S}$ ,  $1200\text{--}1400 \text{ Da}$ ) were also present. This could be elongated monomers, which due to their shape would have higher frictional ratios and underestimated molecular weights.<sup>51</sup> The apparent molecular weight ( $\text{MW}_{\text{app}}$ ) of 4 was much higher than the  $\text{MW}_{\text{ms}}$  of a putative trimer. We conclude that this discrepancy was caused by a tendency of 4 to aggregate. Notably, this made SE AUC analysis impossible, as the peptide aggregated before equilibrium could be attained at any speed. For the other peptides, the trimeric state was confirmed using SE AUC. The data obtained using this technique were fitted to models of single species, monomer-dimer, monomer-trimer, or monomer-tetramer equilibria. The best fits with  $\text{MW}_{\text{app}}$  compliant with  $\text{MW}_{\text{ms}}$  were obtained for the monomer-trimer model for all peptides (Table 1).

The incorporation of *trans*-ACPC did not significantly affect the self-affinity of peptide 1 –  $\log K_a$  of the monomer-trimer equilibria was around 10 for all variants tested. In contrast, substitution with *cis*-ACPC significantly decreased the stability of the obtained peptides compared to model peptide 1. The sedimentation coefficient distributions ( $c(s)$ ) obtained using SV AUC transitioned from more or less regular distributions indicating the presence of trimers to wide irregular distributions indicating the presence of a mixture of loosely bound monomers, dimers, and trimers in the case of 5 and 6, or to a mixture



**Table 1** Comparison of the results of sedimentation velocity (SV AUC) and sedimentation equilibrium (SE AUC) analytical ultracentrifugation. SV AUC was conducted at 20 °C at 50 000 rpm, SE AUC was carried out at 20 °C at 20 000, 35 000 and 48 000 rpm<sup>a</sup>

Peptide	MW <sub>ms</sub> [Da]	$\bar{v}$ [ml g <sup>-1</sup> ]	SV AUC			MW <sub>app</sub> [Da]	<i>n</i>	$\bar{s}_{20,w}$ [S]	<i>ff</i> <sub>0</sub>	SE AUC			
			<i>s</i> [S]	<i>s</i> <sub>20,w</sub> [S]	<i>s</i> <sub>20,w</sub> [S]					Oligomer	log <i>K</i> <sub>a</sub>	MW <sub>app</sub> [Da]	<i>n</i>
1	2713.13	0.76619	0.914	0.950	8182	3.02	0.904	1.314	Trimer	10.0	2814	1.04	
2	2767.22	0.77225	0.913	0.950	8301	3.00	0.902	1.289	Trimer	9.95	2923	1.06	
3	2696.14	0.77439	0.926	0.964	7995	2.97	0.936	1.227	Trimer	10.4	2941	1.09	
4	2821.31	0.77807	0.905	0.943	9395	3.33	0.984	1.373	—	—	—	—	
5	2767.22	0.77225	0.650	0.677	5768	2.08	0.677	1.420	Trimer	8.81	2335	0.85	
6	2696.14	0.77439	0.680	0.708	6254	2.32	0.708	1.419	Trimer	8.57	2655	0.98	
7	2821.31	0.77807	0.422	0.440	3283	1.16	0.481	1.459	Trimer	6.91	3172	1.12	
			0.846	0.881	9315	3.30							

<sup>a</sup> MW<sub>ms</sub> – Molecular Weight determined using mass spectrometry. MW<sub>app</sub> – apparent molecular weight. *n* – quotient of MW<sub>app</sub> and MW<sub>ms</sub>.  $\bar{v}$  – partial specific volume. *s* – sedimentation coefficient. *s*<sub>20,w</sub> – sedimentation coefficient corrected for water, 20 °C.  $\bar{s}_{20,w}$  – weight averaged *s*<sub>20,w</sub>. *ff*<sub>0</sub> – frictional ratio. *K*<sub>a</sub> – association constant.

of monomers and trimers dominated by monomers in the case of 7 (Table 1). The frictional ratio for *cis*-peptides increased from 1.22–1.37 to around 1.45. This indicates that the hydrodynamic drag increased, supposedly as a result of the more elongated conformation of the peptides. SE AUC analyses indicated that the monomer-trimer model is still valid for the *cis*-peptides. The self-affinity was smaller compared to the *trans*-peptides as indicated by the decrease in log *K*<sub>a</sub> from 10 to 8.81 (5) 8.57 (6), and 6.91 (7).

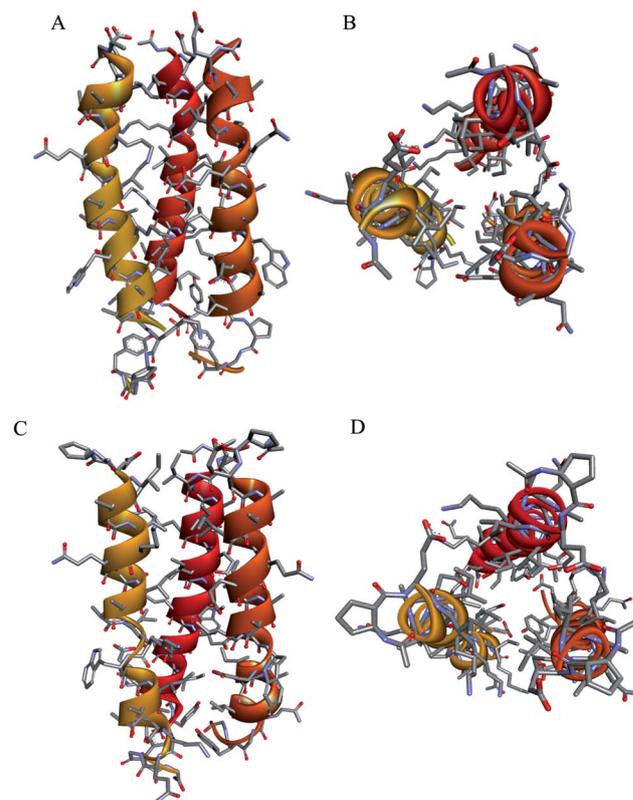
In summary, the AUC results showed that foldamerization did not influence the oligomeric state of the studied peptides. However, substitution with *cis*-ACPC significantly decreased the stability of the obtained peptides compared to model peptide 1, as was also shown by the CD results.

### NMR studies

The peptides with *trans*-ACPC substitution (2 and 3) were chosen for detailed NMR analysis. The unambiguous assignment of nearly all resonances (Tables S2 and S3<sup>†</sup>) was possible based on the analysis of <sup>1</sup>H NMR (Fig. S4 and S7<sup>†</sup> for 2 and 3 respectively), TOCSY (Fig. S5 and S8<sup>†</sup>), and 2D NOESY (Fig. S6 and S9<sup>†</sup>) spectra measured in water solution (pH = 7). In both cases, a significant number of non-sequential inter-proton contacts were identified. Most of the contacts detected were of type *i*–*i* + 3 or *i*–*i* + 4, suggesting the presence of a  $\alpha$ -helical structure consistent with other experiments. Contacts, which were not possible within a helix, were considered to be inter-helical. Based on NMR-derived restraints, the three-dimensional structures of peptides 2 and 3 were calculated (Fig. 4, Table S4<sup>†</sup>).

As expected, peptides 2 and 3 adopt a well-defined coiled-coil conformation in water solution (Fig. S13<sup>†</sup>). Positioning of the isoleucine residue side chains forms a hydrophobic core, while the rest of the side chains are exposed to solvent. Although the *trans*-ACPC residue is placed at the end of the helices, its position is well-defined. Cyclopentane incorporated in *trans*-ACPC does not contribute to the formation of a hydrophobic core as it is directed toward the solvent.

<sup>1</sup>H NMR spectra of *cis*-ACPC mono-substituted peptides 5 and 6 (Fig. S10 and S11<sup>†</sup>) showed a significant decrease in the dispersion of chemical shifts of proton signals and broadening of resonance lines if compared to their *trans*-ACPC-substituted analogues 2 and 3. Two-dimensional TOCSY and NOESY spectra were also recorded for these peptides. However, signal broadening and overlapping disable the precise spectral analysis, suggesting the absence of a well-defined structure, which is clearly visible in the comparison of the representative



**Fig. 4** The averaged structures of peptides 2 and 3 calculated on the basis of restraints derived from 2D NOESY: side view (A and C) and top view (B and D) for 2 and 3, respectively.



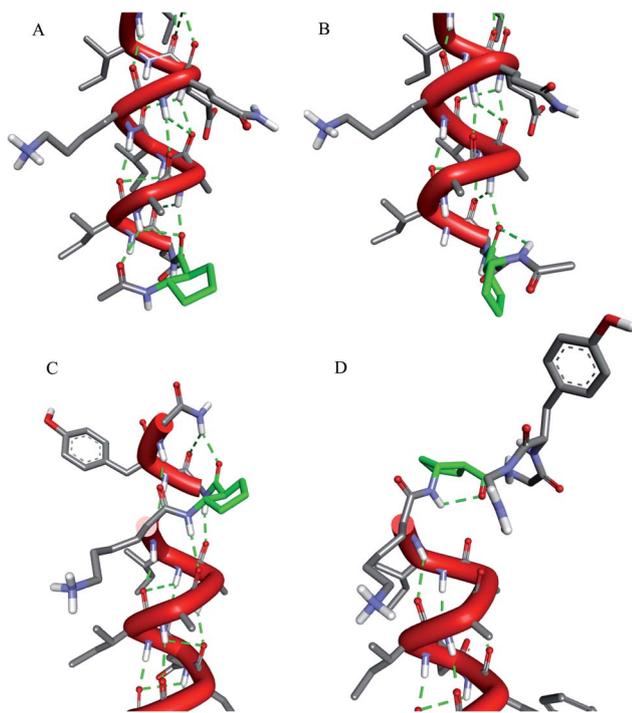


Fig. 5 Zoomed-in fragments of minimized models of peptides containing *trans*-ACPC (sequences 2 and 3, panels A and C, respectively) and *cis*-ACPC (sequences 5 and 6, panels B and D, respectively). ACPC residues are shown in green, and hydrogen bonds are shown as green dotted lines.

fragments of 2D NOESY spectra recorded for *cis*-ACPC (6) and *trans*-ACPC-containing peptides (3) (Fig. S12†). The picture given by NMR measurement is consistent with the CD data and indicates only partial folding of peptides containing the *cis*-ACPC residue.

Moreover, we have modelled the lowest energy conformations of peptides 2, 3, 5 and 6 in order to analyze network of hydrogen bond around  $\beta$ -amino acid residues (Fig. 5). *Trans*-ACPC residue present in the structures of peptides 2 and 3 forms hydrogen bonds analogous to those observed in  $\alpha$ -helix, thus it is stabilizing the helical structure (Fig. 5A and C, respectively). However, the set of hydrogen bonds formed by *cis*-ACPC present in peptides 5 and 6 is different (Fig. 5B and D, respectively). Intra-residue hydrogen bond is observed in both cases, and the network of inter-residue hydrogen bonds is affected. Moreover, the  $\beta$ -amino acid residue, due to its constrained conformation, directs neighboring residues (Ac of peptide 5 and Gly-Tyr of peptide 6) outside the helix.

## Conclusions

Although there are several studies on the effect of substitution of  $\beta$ -amino acids on conformational stability of secondary, tertiary or quaternary structures,<sup>35,48,52</sup> the comparison of analogues containing units of various stereochemistry are rare, and concerns only single  $\alpha$ -helix.<sup>53</sup> Here, we applied well-understood coiled-coil structure as a model for better understanding of effects of *trans*-ACPC and *cis*-ACPC substitutions on

its conformational stability. However, both *trans*-ACPC and *cis*-ACPC can be used as building blocks for construction of various helical structures,<sup>7,45,54</sup> their effect on  $\alpha$ -helix-based coiled-coil is significantly different, as shown in the combined picture derived from all the applied experimental techniques. In particular, CD analysis indicate large difference in conformational stability expressed by  $T_m$  values ranging from approximately 40 °C to 60 °C for *cis*-ACPC and *trans*-ACPC substituted peptides respectively. The difference in conformational stability was also observed in NMR measurements, as only spectra of *trans*-ACPC-based peptides showed numerous non-sequential inter-proton contacts allowing building of consistent model. Moreover, AUC measurements confirmed the trimeric structure of all peptides. Importantly, these detailed structural studies indicate that introduced  $\beta$ -amino acid residues do not interfere with hydrophobic core of the coiled-coil or form any direct inter-helical interactions. Therefore, the striking difference of conformational stability between the corresponding *trans*-ACPC and *cis*-ACPC mutants is a direct result of their ability to promote helical conformation or not. The insight in conformational preferences of these arrangements was achieved by further molecular modelling studies that revealed that *cis*-ACPC intra-residue hydrogen bond (C=O $\cdots$ H-N) degrades the network of hydrogen bonds stabilizing the helix, while *trans*-ACPC adjusts well to  $\alpha$ -helix structure.

It is worth to underline that pairs of peptides 2 and 5, as well as 3 and 6 differ only by one stereogenic center, while their conformational stability are significantly different. Moreover, the discussed changes are introduced at the peripheries of the helices; thus long-range propagation of this effect is observed. Therefore, the small change related to the modulation of the folding propensity of a single amino acid residue has dramatically changed the folding of the coiled-coil structure. In summary, the concept of studying analogues with conformational stability significantly lower than that of the original oligomer can be considered to be complementary to widely studied foldamers and provide a better understanding of the folding properties of various oligomers.

## Experimental section

### Peptide synthesis

All commercially available reagents and solvents were purchased from Lipopharm.pl, Sigma-Aldrich, or Merck and used without further purification. Fmoc-(1*S*, 2*R*)-2-aminocyclopentanecarboxylic acid (Fmoc-*cis*-ACPC) was synthesized according to the known protocol.<sup>55</sup> Fmoc-(1*S*, 2*S*)-2-aminocyclopentanecarboxylic acid (Fmoc-*trans*-ACPC) was purchased from Synnovator, Inc. The peptides described were obtained with an automated solid-phase peptide synthesizer (Biotage® Initiator + Alastra™) using rink amide AM resin (loading: 0.59 mmol g<sup>-1</sup>). Fmoc deprotection was performed using 20% piperidine in DMF for 3 + 10 min. A double coupling procedure was achieved with 0.5 M DIC solution and 0.5 M OXYMA solution (1 : 1) in DMF, for  $\alpha$ -amino acids 2  $\times$  15 min and for  $\beta$ -amino acids 30 min, at 75 °C. The acetylation reaction was carried out using a mixture of NMP/DIPEA/acetic anhydride



(80 : 15 : 5). The cleavage of the peptides from the resin was accomplished with the mixture of TFA/TIS/H<sub>2</sub>O (95 : 2.5 : 2.5) after 3 h of shaking. The crude peptide was precipitated with ice-cold Et<sub>2</sub>O and centrifuged (10 000 rpm, 15 min, 4 °C). The peptides were purified using preparative HPLC (Knauer Prep) with a C18 column (Thermo Scientific, Hypersil Gold 12 μ, 250 mm × 20 mm) with a water/acetonitrile (0.05% TFA) eluent system.

### Analytical high-performance liquid chromatography (HPLC)

Analytical HPLC was performed using a Kinetex 5μ EVO C18 100A 150 × 4.6 mm column. Program (eluent A: 0.05% TFA in H<sub>2</sub>O, eluent B: 0.05% TFA in acetonitrile, flow 0.5 mL min<sup>-1</sup>): A: *t* = 0 min, 90% A; *t* = 30 min, 10% A.

### Mass spectrometry (MS)

Mass spectra were recorded using a WATERS LCT Premier XE high-resolution mass spectrometer with an electrospray ionization and time-of-flight (TOF) detector.

### Circular dichroism (CD)

CD spectra were recorded using a JASCO J-815 spectropolarimeter at 20 °C between 250 and 190 nm in water with the following parameters: 0.2 nm resolution, 1.0 nm bandwidth, 20 mdeg sensitivity, 0.25 s response, 50 nm min<sup>-1</sup> scanning speed, 5 accumulations, 0.02 cm cuvette path length. The CD spectra of the buffer alone were recorded and subtracted from the raw data. Typically, samples were prepared by dilution of a peptide stock solution at a concentration of 1.3 mM. CD intensity is given as the mean residue molar ellipticity ( $\theta$  [deg × cm<sup>2</sup> × dmol<sup>-1</sup>]). The melting temperatures of the studied peptides were determined by a method described previously.<sup>56</sup>

### Temperature denaturation measurements using CD

To examine the thermal unfolding of the peptide, stock solutions were diluted to 80 μM and two types of measurements were performed: (1) in the range of 190–240 nm and (2) at 222 nm in phosphate buffer pH = 7, 0.05 M. The temperature increased from 4 to 98 °C in increments of 2 °C. Ellipticity measurements were recorded with a 1 mm path length cuvette and other parameters remained unchanged.

### Analytical ultracentrifugation (AUC)

The weighed portions of the peptides were dissolved in PBS (7.74 mM Na<sub>2</sub>HPO<sub>4</sub>, 2.26 mM NaH<sub>2</sub>PO<sub>4</sub>, 137 mM NaCl, 2.7 mM NaCl, pH 7.4) at room temperature for at least 15 minutes in an Eppendorf Thermomixer at 500 rpm and subsequently centrifuged at 14 000g for 5 minutes at room temperature. The supernatants were collected and the concentrations of the peptides were calculated from absorbance values measured at 280 nm (*A*<sub>280</sub>) and molar extinction coefficients of 6990 M<sup>-1</sup> cm<sup>-1</sup>. Sedimentation velocity (SV AUC) and sedimentation equilibrium (SE AUC) analytical ultracentrifugation experiments were conducted at 20 °C using a Beckman Coulter ProteomeLab XLI analytical ultracentrifuge and an An-60Ti rotor. Partial specific volumes of

the peptides were estimated using SEDNTERP<sup>57</sup> (available at <http://www.jphilo.mailway.com/download.htm>, 6th April 2020) and corrected for the presence of ACPC, N-terminal acetyl and C-terminal amine moieties using published molar increment values of chemical groups.<sup>58</sup> Density and viscosity of PBS were estimated using SEDNTERP. SV AUC was conducted in cells containing charcoal-filled Epon centerpieces with 2 sector-shaped channels for 400 μL of 100 μM peptides in PBS. The samples were centrifuged overnight (approximately 20 h) at 50 000 rpm with continuous collection of A280 with 0.003 cm resolution. The time-corrected data<sup>59</sup> were fitted to a continuous sedimentation coefficient distribution (*c*(*s*)) model<sup>60</sup> using SEDFIT software (available at <https://sedfitsedphat.nibib.nih.gov/software/default.aspx>, 6th April 2020). The fits of 150 sedimentation coefficient values between 0 and 3 S were calculated alternately using simplex and Marquardt–Levenberg algorithms. The quality of the fits was estimated using RMSD values. A *p*-value for regularization was set at 0.95. SE AUC was conducted at 20 000, 35 000 and 48 000 rpm in cells containing charcoal-filled Epon centerpieces with 6 square channels for 100 μL of approximately 50 μM peptides. For peptide 7, an additional experiment was conducted for 50, 75 and 100 μM peptide samples at 38 000, 41 000, and 45 000 rpm to better describe its properties. The scans of intensities of light at 280 nm were collected separately for reference and sample channels in step mode with 0.001 cm resolution and 10 replicates per scan. Scans representing the peptide solutions at equilibrium were processed using GUSI (available at <http://biophysics.swmed.edu/MBR/software.html>, 6th April 2020)<sup>61</sup> and analyzed using SEDPHAT (available at <https://sedfitsedphat.nibib.nih.gov/software/default.aspx>, 6th April 2020) using single species, monomer-dimer, monomer-trimer and monomer-tetramer equilibrium models. The fits were calculated alternately using simplex and Marquardt–Levenberg algorithms. The final fit was performed using a simulated annealing algorithm with default settings. The accuracy of the fits was followed using global RMSD values.

### Nuclear magnetic resonance (NMR)

**Nuclear magnetic resonance.** NMR experiments were performed on a Bruker Avance™ spectrometer operating at 600.58 MHz for <sup>1</sup>H, equipped with a 5.0 mm PA BBO probe. The NMR spectra at temperature 298 K were recorded in 10% D<sub>2</sub>O/water and adjusted to pH 7 by NaOH solution. The temperature was controlled to ±0.1 K. TOCSY and NOESY experiments were performed for chemical shift and structure assignment. All NMR spectra were acquired with suppression of the solvent OH signal using a 3-9-19 pulse sequence with gradients. Typical TOCSY – homonuclear Hartman (Hahn transfer using the mlev17 sequence for mixing using two power levels for excitation and spinlock) and 2D NOESY were recorded in phase-sensitive mode with the spectral width of 6127 Hz in both dimensions using 2048 data points and relaxation delay of 2 s. These spectra were acquired with 1024 increments of 16 scans for the TOCSY and 80 scans for 2D NOESY. Mixing times were set at 80 ms and 250 ms for the TOCSY and 2D NOESY experiments, respectively. Data were acquired and processed using



Topspin 3.1 (BrukerBioSpin, Rheinstetten, Germany). The processed spectra were assigned with the help of the SPARKY program.<sup>62</sup>

**NMR structure calculation.** The NMR structure generation was performed in Xplor-NIH v. 2.41.1 program.<sup>63</sup> Initially, 100 random conformations were generated from the model structure of a trimer containing three identical chains using the pdbTOPSF protocol. NMR-derived interproton contacts were classified by standard method with upper distance limits: strong 2.5 Å, medium 3.5 Å and weak 5 Å, and the lower distance limit was set to 1.8 Å. The contacts between the protons belonging to the side chains of vicinal residues that are not likely for the helix structure were used as interhelical distance restraints, and the rest contacts were used as intrahelical distance restraints. For the  $\beta$ -amino acid also 2 backbone torsions were restraint to match the previous experimental data<sup>64</sup> with a tolerance of 30 degree deviation. Standard simulated annealing protocols implemented in Xplor-NIH were used composed of the following steps: (1) high temperature dynamics (3500 K, 800 ps or 8000 steps), (2) simulated annealing performed from 3500 K to 25 K with 12.5 K step, at each temperature short dynamics was done (100 steps or 0.2 ps); (3) gradient minimization of final structure. Finally, the top ten lowest energy structures were superimposed and averaged.

**Molecular modelling.** Models of peptides 2, 3, 5 and 6 were prepared using Discovery Studio Visualiser v20 on the basis of the experimental structure of a *de novo* designed coiled-coil (pdb: 4dzk).<sup>40</sup> The starting *trans*-ACPC conformation in peptides 2 and 3 was taken from the crystal structure of the peptide containing *trans*-ACPC residues that mimics  $\alpha$ -helical structure (pdb: 3f50),<sup>7</sup> while the *cis*-ACPC conformation in peptides 5 and 6 was applied from the crystal structure of the helical peptide containing *cis*-ACPC at a terminal position (pdb: 7ad0).<sup>53</sup> The prepared models were then submitted to minimization using the GROMACS software (available on WCSS, Wrocław, Poland) applying the Amber03 force field extended with noncanonical amino acids.<sup>65</sup> The box definition and the solvation of the system were performed using gmx editconf and gmx solvate methods with the spc216 water model. The minimization of the system was performed applying the steepest descent algorithm with 5000 maximum number of steps and PME electrostatic until  $F_{\max} < 1000 \text{ kJ mol}^{-1} \text{ nm}^{-1}$ . For every peptide the model converged in less than 400 steps. Minimization was followed by two-phase equilibration: 100 ps (50 000 steps) under the canonical ensemble (NVT) and 100 ps (50 000 steps) under the NPT ensemble.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

The authors would like to gratefully acknowledge National Science Centre, Poland for their financial support through grant no. 2017/26/D/ST5/00341. The Wrocław Center for Networking and Supercomputing (WCSS) is kindly acknowledged for the use

of software (GROMACS) and hardware resources (computing grant of Ł. B.).

## References

- 1 S. H. Gellman, *Acc. Chem. Res.*, 1998, **31**, 173–180.
- 2 A. N. Lupas and J. Bassler, *Trends Biochem. Sci.*, 2017, **42**, 130–140.
- 3 E. H. C. Bromley and K. J. Channon, *Prog. Mol. Biol. Transl. Sci.*, 2011, **103**, 231–275.
- 4 D. N. Woolfson, *Adv. Protein Chem.*, 2005, **70**, 79–112.
- 5 W. S. Horne, J. L. Price, J. L. Keck and S. H. Gellman, *JACS*, 2007, **129**, 4178–4180.
- 6 W. S. Horne, M. D. Boersma, M. A. Windsor and S. H. Gellman, *Angew. Chem., Int. Ed.*, 2008, **47**, 2853–2856.
- 7 W. S. Horne, L. M. Johnson, T. J. Ketas, P. J. Klasse, M. Lu, J. P. Moore and S. H. Gellman, *Proc. Natl. Acad. Sci. U. S. A.*, 2009, **106**, 14751–14756.
- 8 L. K. A. Pilsl and O. Reiser, *Amino Acids*, 2011, **41**, 709–718.
- 9 W. S. Horne and S. H. Gellman, *Acc. Chem. Res.*, 2008, **41**, 1399–1408.
- 10 T. A. Martinek and F. Fulop, *Chem. Soc. Rev.*, 2012, **41**, 687–702.
- 11 C. M. Goodman, S. Choi, S. Shandler and W. F. DeGrado, *Nat. Chem. Biol.*, 2007, **3**, 252–362.
- 12 P. G. Vasudev, S. Chatterjee, N. Shamala and P. Balaram, *Chem. Rev.*, 2011, **111**, 657–687.
- 13 Y.-H. Shin and S. H. Gellman, *JACS*, 2018, **140**, 1394–1400.
- 14 S. Rinaldi, *Molecules*, 2020, **25**, 3276.
- 15 J. Briocche, S. J. Pike, S. Tshepelevitsh, I. Leito, G. A. Morris, S. J. Webb and J. Clayden, *J. Am. Chem. Soc.*, 2015, **137**, 6680–6691.
- 16 L. Byrne, J. Sola and J. Clayden, *Chem. Commun.*, 2015, **51**, 10965–10968.
- 17 M. De Poli, W. Zawodny, O. Quinonero, M. Lorch, S. J. Webb and J. Clayden, *Science*, 2016, **352**, 575–580.
- 18 G. W. Collie, K. Pulka-Ziach, C. M. Lombardo, J. Fremaux, F. Rosu, M. Decossas, L. Mauran, O. Lambert, V. Gabelica, C. D. Mackereth and G. Guichard, *Nat. Chem.*, 2015, **7**, 871–878.
- 19 R. Chingle, C. Proulx and W. D. Lubell, *Acc. Chem. Res.*, 2017, **50**, 1541–1556.
- 20 P. Amabili, M. Calvaresi, G. Martelli, M. Orena, S. Rinaldi and F. Sgolastra, *Eur. J. Org. Chem.*, 2019, **2019**, 907–917.
- 21 A. Lobo-Ruiz and J. Tulla-Puche, 2 – Synthetic approaches of naturally and rationally designed peptides and peptidomimetics, *Peptide Applications in Biomedicine, Biotechnology and Bioengineering*, ed. S. Koutsopoulos, Woodhead Publishing, 2018, pp. 23–49.
- 22 J. Vagner, H. Qu and V. J. Hruby, *Curr. Opin. Chem. Biol.*, 2008, **12**, 292–296.
- 23 E. Lenci and A. Trabocchi, *Chem. Soc. Rev.*, 2020, **49**, 3262–3277.
- 24 M. Muttenthaler, G. F. King, D. J. Adams and P. F. Alewood, *Nat. Rev. Drug Discovery*, 2021, **20**, 309–325.
- 25 G. Olajos, A. Hetenyi, E. Weber, T. Szogi, L. Fulop and T. A. Martinek, *Org. Biomol. Chem.*, 2018, **16**, 5492–5499.



- 26 D. Seebach, A. K. Beck and D. J. Bierbaum, *Chem. Biodiversity*, 2004, **1**, 1111–1239.
- 27 A. I. Webb, M. A. Dunstone, N. A. Williamson, J. D. Price, A. de Kauwe, W. Chen, A. Oakley, P. Perlmutter, J. McCluskey, M.-I. Aguilar, J. Rossjohn and A. W. Purcell, *J. Immunol.*, 2005, **175**, 3810–3818.
- 28 P. Zubrzak, H. Williams, G. M. Coast, R. E. Isaac, G. Reyes-Rangel, E. Juaristi, J. Zabrocki and R. J. Nachman, *Biopolymers*, 2007, **88**, 76–82.
- 29 S. Reinelt, M. Marti, S. Dedier, T. Reitingner, G. Folkers, J. A. Lopez de Castro and D. Rognan, *J. Biol. Chem.*, 2001, **276**, 24525–24530.
- 30 J. D. Sadowsky, J. K. Murray, Y. Tomita and S. H. Gellman, *ChemBioChem*, 2007, **8**, 903–916.
- 31 L. Berlicki, M. Kaske, R. Gutierrez-Abad, G. Bernhardt, O. Illa, R. M. Ortuno, C. Cabrele, A. Buschauer and O. Reiser, *Med. Chem.*, 2013, **56**, 8422–8431.
- 32 U. Strijowski and N. Sewald, *Org. Biomol. Chem.*, 2004, **2**, 1105–1109.
- 33 E. Vass, U. Strijowski, K. Wollschlager, I. M. Mandity, G. Szilvagy, M. Jewginski, K. Gaus, S. Royo, Z. Majer, N. Sewald and M. Hollosi, *J. Pept. Sci.*, 2010, **16**, 613–620.
- 34 D. Seebach, A. Lukaszuk, K. Patora-Komisarska, D. Podwysocka, J. Gardiner, M.-O. Ebert, J. C. Reubi, R. Cescato, B. Waser, P. Gmeiner, H. Hubner and C. Rougeot, *Chem. Biodiversity*, 2011, **8**, 711–739.
- 35 M. Szeferczyk, N. Szulc, M. Gasior-Głogowska, A. Modrak-Wojcik, A. Bzowska, W. Majstrzyk, M. Taube, M. Kozak, T. Gotszalk, E. Rudzinska-Szostak and L. Berlicki, *Nanoscale*, 2021, **13**, 4000–4015.
- 36 M. Oba, C. Ito and M. Tanaka, *Bioorg. Med. Chem. Lett.*, 2018, **28**, 875–877.
- 37 M. Drewniak-Switalska, B. Barycza, E. Rudzinska-Szostak, P. Morawiak and L. Berlicki, *Org. Biomol. Chem.*, 2021, **19**, 4272–4278.
- 38 F. Lapenta, J. Aupic, Z. Strmsek and R. Jerala, *Chem. Soc. Rev.*, 2018, **47**, 3530–3542.
- 39 P. B. Harbury, T. Zhang, P. S. Kim and T. A. Alber, *Science*, 1993, **262**, 1401–1407.
- 40 J. M. Fletcher, A. L. Boyle, M. Bruning, G. J. Bartlett, T. L. Vincent, N. R. Zaccai, C. T. Armstrong, E. H. C. Bromley, P. J. Booth, R. L. Brady, A. R. Thomson and D. N. Woolfson, *ACS Synth. Biol.*, 2012, **1**, 240–250.
- 41 W. S. Horne, J. L. Price and S. H. Gellman, *Proc. Natl. Acad. Sci. U. S. A.*, 2008, **105**, 9151–9156.
- 42 J. L. Price, E. B. Hadley, J. D. Steinkruger and S. H. Gellman, *Angew. Chem., Int. Ed.*, 2010, **49**, 368–371.
- 43 B. F. Fisher, S. H. Hong and S. H. Gellman, *JACS*, 2018, **140**, 9396–9399.
- 44 I. M. Mandity, E. Weber, T. A. Martinek, G. Olajos, G. K. Toth, E. Vass and F. Fulop, *Angew. Chem., Int. Ed.*, 2009, **48**, 2171–2175.
- 45 L. Berlicki, L. Pils, E. Weber, I. M. Mandity, C. Cabrele, T. A. Martinek, F. Fulop and O. Reiser, *Angew. Chem., Int. Ed.*, 2012, **51**, 2208–2212.
- 46 M. Szeferczyk, E. Weglarz-Tomczak, P. Fortuna, A. Krzyszton, E. Rudzinska-Szostak and L. Berlicki, *Angew. Chem., Int. Ed.*, 2017, **56**, 2087–2091.
- 47 J. L. Price, W. S. Horne and S. H. Gellman, *JACS*, 2010, **132**, 12378–12387.
- 48 D. E. Mortenson, D. F. Kreitler, N. C. Thomas, I. A. Guzei, S. H. Gellman and K. T. Forest, *ChemBioChem*, 2018, **19**, 604–612.
- 49 C. Toniolo, A. Polese, F. Formaggio, M. Crisma and J. Kamphuis, *JACS*, 1996, **118**, 2744–2745.
- 50 A. Micsonai, F. Wien, L. Kernya, Y.-H. Lee, Y. Goto, M. Refregiers and J. Kardos, *Proc. Natl. Acad. Sci. U. S. A.*, 2015, **112**, E3095–E3103.
- 51 P. Schuck, H. Zhao, C. A. Brautigam and R. Ghirlando, *Basic Principles of Analytical Ultracentrifugation*, Boca Raton, FL, CRC Press, 2015.
- 52 M. Drewniak, E. Weglarz-Tomczak, K. Ozga, E. Rudzinska-Szostak, K. Macegoniuk, J. Tomczak, M. Bejger, W. Rypniewski and L. Berlicki, *Bioorg. Chem.*, 2018, **81**, 356–361.
- 53 P. Fortuna, A. Twarda-Clapa, Ł. Skalniak, K. Ożga, T. A. Holak and Ł. Berlicki, *Eur. J. Med. Chem.*, 2020, **208**, 112814.
- 54 T. A. Martinek, A. Hetenyi, L. Fulop, I. M. Mandity, G. K. Toth, I. Dekany and F. Fulop, *Angew. Chem., Int. Ed.*, 2006, **45**, 2396–2400.
- 55 S. G. Davies, O. Ichihara, I. Lenoir and I. A. S. Walters, *J. Chem. Soc., Perkin Trans. 1*, 1994, **11**, 1411–1415.
- 56 D. F. Kreitler, D. E. Mortenson, K. T. Forest and S. H. Gellman, *JACS*, 2016, **138**, 6498–6505.
- 57 T. M. Laue, B. D. Shah, T. M. Ridgeway and S. L. Pelletier, *Analytical Ultracentrifugation in Biochemistry and Polymer Science*, Royal Society of Chemistry, ed. S. Harding and A. Rowe, 1992, pp. 90–125.
- 58 H. Durchschlag and P. Zipper, *J. Appl. Crystallogr.*, 1997, **30**, 803–807.
- 59 H. Zhao, R. Ghirlando, G. Piszczek, U. Curth, C. A. Brautigam and P. Schuck, *Anal. Biochem.*, 2013, **437**, 104–108.
- 60 P. Schuck, *Biophys. J.*, 2000, **78**, 1606–1619.
- 61 C. A. Brautigam, *Methods Enzymol.*, 2015, **562**, 109–133.
- 62 T. D. Goddard and D. G. Kneller, *Sparky, 3rd edn*, University of California, San Francisco, CA, USA, 2001.
- 63 C. D. Schwieters, J. J. Kuszewski and G. M. Clore, *Prog. Nucl. Magn. Reson. Spectrosc.*, 2006, **48**, 47–62.
- 64 T. A. Martinek, G. K. Toth, E. Vass, M. Hollosi and F. Fulop, *Angew. Chem., Int. Ed.*, 2002, **41**, 1718–1721.
- 65 L. J. Németh, Z. Hegedüs and T. A. Martinek, *J. Chem. Inf. Model.*, 2014, **54**, 2776–2783.

