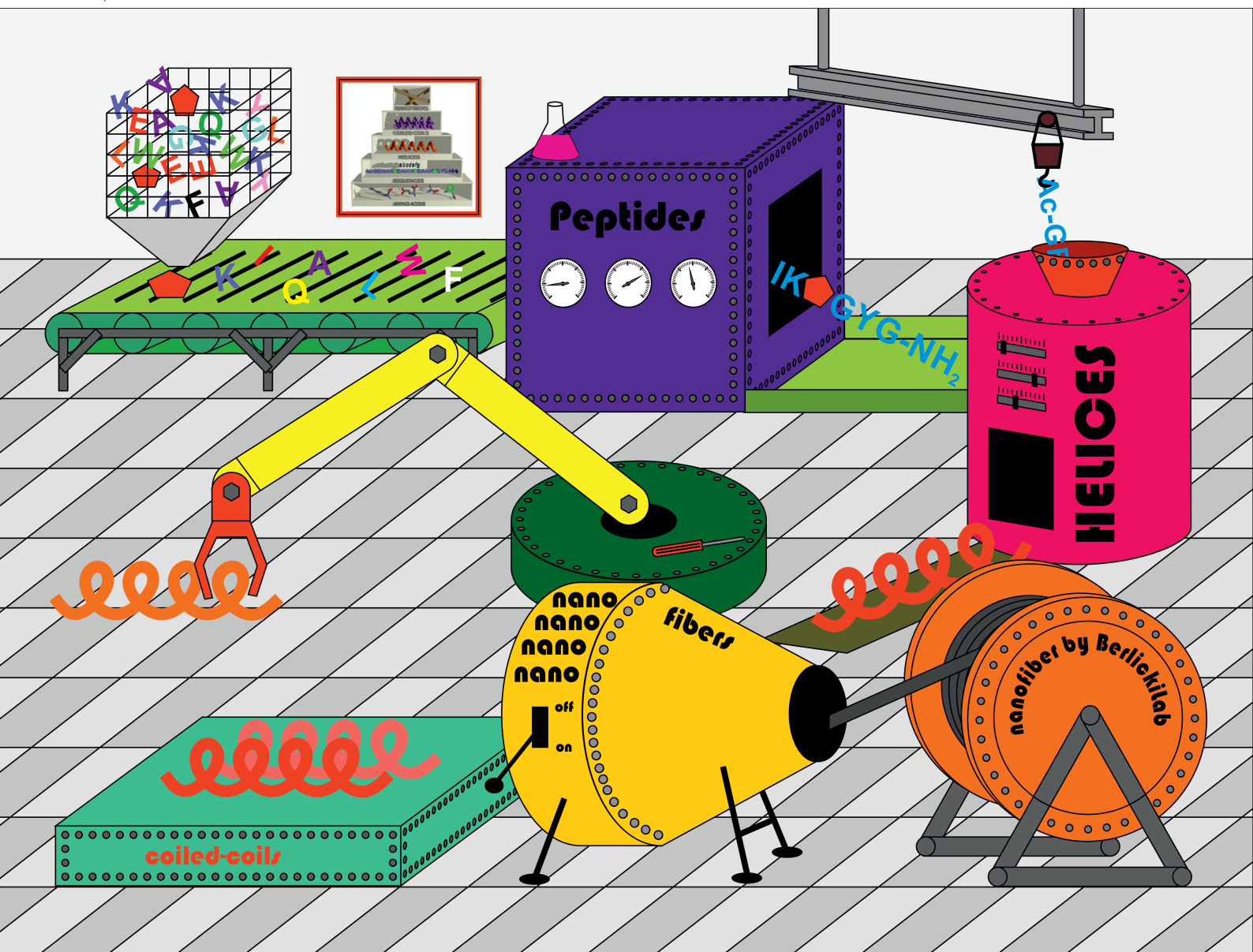


Nanoscale

rsc.li/nanoscale



ISSN 2040-3372



Cite this: *Nanoscale*, 2021, **13**, 4000

Hierarchical approach for the rational construction of helix-containing nanofibrils using α,β -peptides†

Monika Szeftczyk, *^a Natalia Szulc, ^b Marlena Gąsior-Głogowska, ^b Anna Modrak-Wójcik, ^c Agnieszka Bzowska, ^c Wojciech Majstrzyk, ^d Michał Taube, ^e Maciej Kozak, ^{e,f} Teodor Gotszalk, ^d Ewa Rudzińska-Szostak ^a and Łukasz Berlicki *^a

The rational design of novel self-assembled nanomaterials based on peptides remains a great challenge in modern chemistry. A hierarchical approach for the construction of nanofibrils based on α,β -peptide foldamers is proposed. The incorporation of a helix-promoting *trans*-(1*S*,2*S*)-2-aminocyclopentanecarboxylic acid residue in the outer positions of the model coiled-coil peptide led to its increased conformational stability, which was established consistently by the results of CD, NMR and FT-IR spectroscopy. The designed oligomerization state in the solution of the studied peptides was confirmed using analytical ultracentrifugation. Moreover, the cyclopentane side chain allowed additional interactions between coiled-coil-like structures to direct the self-assembly process towards the formation of well-defined nanofibrils, as observed using AFM and TEM techniques.

Received 6th June 2020,
Accepted 4th January 2021

DOI: 10.1039/d0nr04313c

rsc.li/nanoscale

Introduction

The rational design of self-assembled nanostructures based on peptides has recently attracted much attention.^{1,2} In this context, helical structures possess numerous advantages, including rational design, high conformational stability and the possibility of a wide range of modifications that could lead to the formation of higher-order structures with different complexities and functions.^{3,4} A coiled-coil is one of the most widely studied helix-containing peptide motifs because of the well-understood relationship between its sequence and structure,^{5–7} which allows the construction of several nanostructures, *i.e.*, bundles,⁸ nanofibers,⁹ nanotubes¹⁰ or cages.¹¹ Fibrous nanostructures based on coiled-coil peptides have

been most often obtained by the formation of salt bridges, disulfide bonds or complementary acidic/basic components.¹² However, the construction of coiled-coil-based nanofibers exploiting hydrophobic interactions between constituent elements remains virtually unexplored.

Rational and effective control over the three-dimensional structures of peptides can be achieved by incorporating conformationally constrained amino acid residues. Among numerous approaches, the application of cycloalkane-based β -amino acids has already been proven to be very successful, and several highly stable helical conformations have been found for peptides containing cyclopropane, cyclobutane, cyclopentane and cyclohexane β -amino acids.^{13–15} Furthermore, peptides containing β -amino acid residues have been used for the construction of supramolecular self-assemblies, including bundles and coiled-coils.^{15–20} Such peptides have also been proven to self-associate to form various structures including: hydrogels stable for proteolysis,^{21,22} liquid crystals²³ or the vitamin B derivatives releasing nanoparticles.²⁴

The methodology for the rational construction of secondary structures based on α,β -peptides, in particular helices, has already been elaborated^{25,26} and is based on the tendency of individual residues to form ψ and ϕ dihedral angles. Stereochemical patterning indicates that a helix will be obtained if the ψ and ϕ dihedral angles flanking the amide bond are of the same sign.²⁷ However, rational methodologies for the construction of higher-order structures or assemblies using α,β -peptides are scarce. The most commonly used approach is based on $\alpha \rightarrow \beta$ replacement, assuming that the incorporation

^aDepartment of Bioorganic Chemistry, Faculty of Chemistry, Wrocław University of Science and Technology, Wybrzeże Wyspiańskiego 27, 50-370 Wrocław, Poland. E-mail: monika.szeftczyk@pwr.edu.pl, lukasz.berlicki@pwr.edu.pl

^bDepartment of Biomedical Engineering, Faculty of Fundamental Problems of Technology, Wrocław University of Science and Technology, Wybrzeże Wyspiańskiego 27, 50-370 Wrocław, Poland

^cDivision of Biophysics, Faculty of Physics, Institute of Experimental Physics, University of Warsaw, Ludwika Pasteura 5, 02-093 Warsaw, Poland

^dFaculty of Microsystem Electronics and Photonics, Wrocław University of Science and Technology, Wybrzeże Wyspiańskiego 27, 50-370 Wrocław, Poland

^eDepartment of Macromolecular Physics, Adam Mickiewicz University, Uniwersytetu Poznańskiego 2, 61-614 Poznań, Poland

^fNational Synchrotron Radiation Centre SOLARIS, Jagiellonian University, Czerwone Maki 98, 30-392 Kraków, Poland

† Electronic supplementary information (ESI) available: Additional figures and tables. See DOI: 10.1039/d0nr04313c



the solid state will be evaluated. We presumed that *trans*-ACPC residues present in the outer positions (*b*, *c* and *f*) of the coiled-coil will provide the additional possibility of interactions between coiled-coil units to favor the formation of larger, nanometer-size structures.

Synthesis and structure elucidation

All designed peptides were successfully synthesized using an automated peptide synthesizer with microwaves by applying Fmoc chemistry and purified using a preparative HPLC system with a C18 column. Initially, circular dichroism (CD) spectroscopy was used to elucidate the secondary structure and conformational stability of the obtained peptides. The CD spectrum of model peptide **1** shows two minima, at 208 and 222 nm, typical for the α -helix (Fig. 2A). The ratio molar ellipticities at the abovementioned wavelengths of $\theta_{222}/\theta_{208}$ equals 1, and θ_{222} equals *ca.* $-40\,000\text{ deg cm}^2\text{ dmol}^{-1}$, indicating a near-complete α -helix structure.³⁶ The obtained α,β -peptides show two different types of conformational behavior, which depend on the placement of cyclic β residues in the sequence. The CD spectra of peptides **1_b**, **1_c** and **1_f** (dissolved in water, pH 7) have two minima, at 208 and 222 nm, but $\theta_{222}/\theta_{208} < 1$, equal to 0.4, 0.7 and 0.7, respectively, indicating the presence of structures similar to the α -helix but disturbed by the presence of the β residue (Fig. 2A). In the case of peptides **1_c** and **1_f**, the CD spectra show a significant Cotton effect, which suggests the formation of well-defined secondary structures. Very similar CD spectra were also observed for peptides dissolved in methanol (Fig. S3_1†). The CD spectra of peptides **1_a**, **1_d**, **1_e** and **1_g** show a minimum in the 204–205 nm range (Fig. 2A), which indicates a random coil.

The conformational stability was examined for helical peptides, namely, **1**, **1_b**, **1_c** and **1_f**. The changes in CD spectra were followed in the range of 190–240 nm by a gradual increase in temperature from 4 to 98 °C (Fig. S3_1†). Thermal unfolding curves followed at 222 nm (Fig. 2B) were used to determine the melting temperatures (Table S3_2†). As

expected, the short coiled-coil peptide **1** has moderate thermal stability in water with a cooperative melting transition at 56 °C. The obtained α,β -peptides also demonstrate cooperative melting transitions but at significantly higher temperatures, 76 °C and 71 °C for **1_c** and **1_f**, respectively. Peptide **1_b** strongly aggregates under the measurement conditions (increase in the temperature), so its melting temperature was not possible to determine. The conformational stability of the peptides was additionally confirmed in the presence of 3 M guanidine hydrochloride, which was used as a denaturing agent, and it was observed that the obtained α,β -peptides exhibit higher conformational stability than peptide **1** under these conditions (Fig. S3_3†). In summary, the results of CD spectroscopy indicate the formation of well-ordered structures for peptides **1_c** and **1_f** that exceed the stability of model short coiled-coil peptide **1**. Moreover, peptide **1_b** shows a partially folded helical structure. Importantly, these results correlate well with the *trans*-ACPC substitution pattern. Only peptides with *trans*-ACPC placed in position designed to be external in the coiled-coil show the formation of a helix. The substitution of positions *a*, *d*, *e* and *g* that form hydrophobic cores or surface hydrogen bond networks leads to complete destruction of the helix.

To elucidate the three-dimensional structure in solution of peptides **1_b**, **1_c** and **1_f**, detailed NMR analysis applying 2D TOCSY and 2D ROESY spectra was performed. Due to the insufficient solubility in water, the measurements for α,β -peptides were performed in methanol-*d*₃ solution. Spectral analysis allowed the assignment of all resonances in all measured peptides and the detection of numerous nonsequential contacts between residues (Table S4†), namely, $i \rightarrow (i - 2)$, $i \rightarrow (i - 3)$ and $i \rightarrow (i - 4)$, which indicate the presence of stable helical structures. In the 2D ROESY spectra of all studied peptides, the most abundant are medium range $i \rightarrow (i - 3)$ contacts (HA-HN, HA-HA and HA-HB). Moreover, peptide **1_f** exhibited a significantly higher number of $i \rightarrow (i - 3)$ nonsequential interproton contacts than two other peptides (36 *versus* 22 in **1_c** and 19 in **1_b**). Using NMR restraints

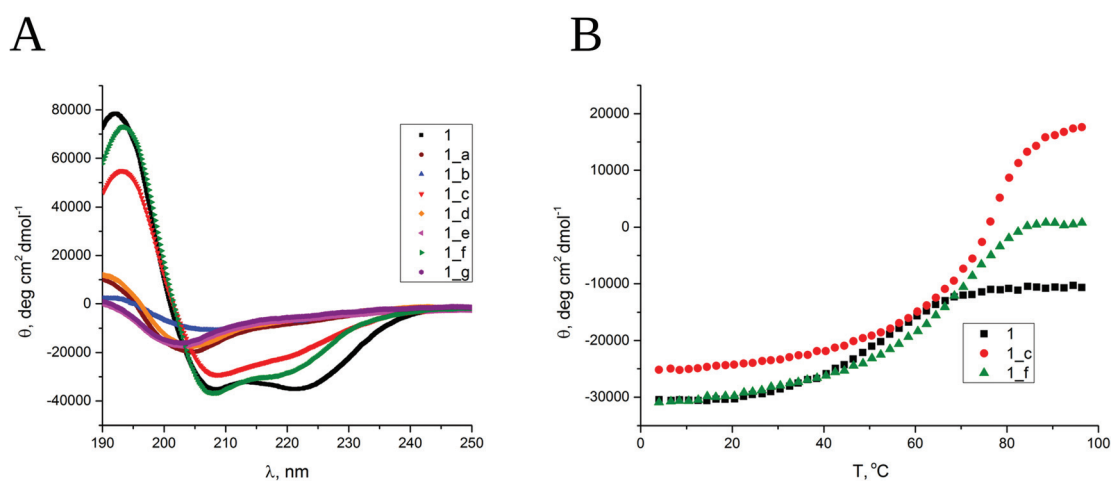


Fig. 2 (A) CD spectra of all obtained peptides dissolved in water, pH 7. (B) Thermal unfolding curves followed by CD signal at 208 nm. $C_{\text{pep}} = 80\ \mu\text{M}$.



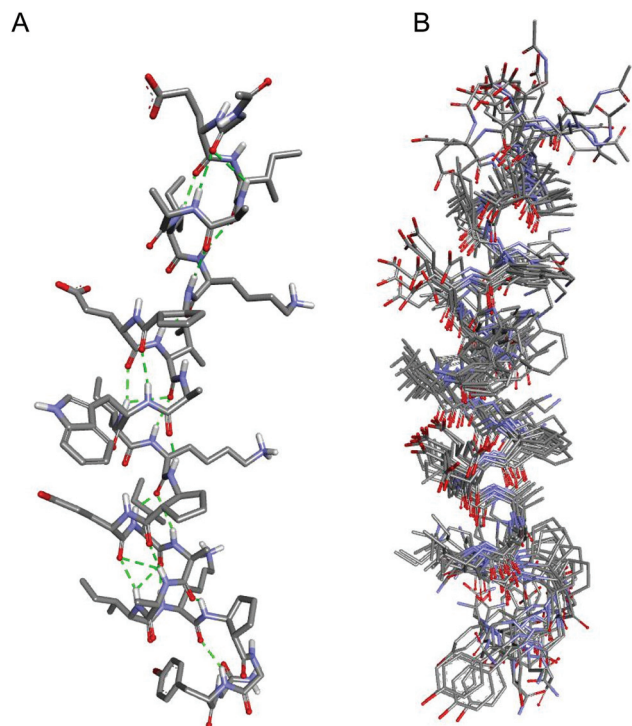


Fig. 3 (A) Average NMR structure and (B) superimposition of the ten lowest energy structures obtained for peptide **1_f**.

(interproton distances and J coupling constants), the three-dimensional structures of peptides **1_b**, **1_c** and **1_f** were modeled (Fig. 3 and S4₁†). As expected, the peptide **1_f** is fully helical, with a structure highly similar to α -helix stabilized by the network of $i \rightarrow (i - 4)$ hydrogen bonds. Although the overall structure is slightly bent due to the disturbances observed at the N-terminal side of each *trans*-ACPC residue, where some hydrogen bonds are elongated or broken, the whole structure is very well defined. Structures of peptide **1_b** and **1_c** are also helical, although their C-terminal part is less defined than in case of peptide **1_f** (Fig. S4₁†). Moreover, the structure of peptide **1_f** trimer was modelled (Fig. S4₂†) and it was found that overall structure is highly similar to native coiled coil with small changes in conformation close to β -amino acid residues.

Aggregate formation

Analytical ultracentrifugation (AUC) was used to verify the oligomerization state of the obtained peptides in solution. For all peptides, sedimentation velocity experiments were carried out in water and in AUC buffer (see Experimental section), except for **1_f**, which strongly aggregates in the buffer. For peptide **1**, sedimentation equilibrium experiments in AUC buffer were also conducted. The $c(s)$ distributions, resulting from the analysis of radial absorption profiles for peptide **1** (Fig. S5₁†), contain a single peak at 0.85–0.97 S slightly depending on the concentration and solvent (s -values are lower in water; Fig. 4 and Table 1). The value of the sedimentation coefficient corres-

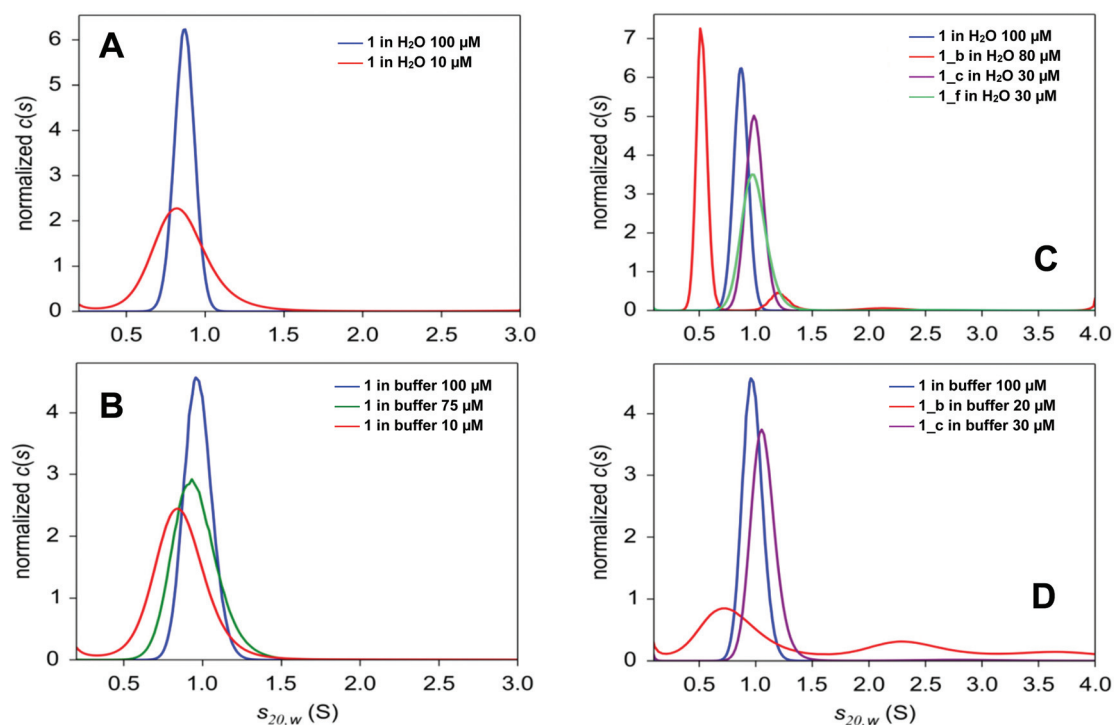


Fig. 4 Normalized sedimentation coefficient distributions $c(s_{20,w})$ for different concentrations of peptide **1** obtained in water (A), in 50 mM phosphate pH 7.0 and 50 mM NaCl (B), as well as for peptides **1**, **1_b**, **1_c** and **1_f** in water (C), in 50 mM phosphate pH 7.0 and 50 mM NaCl (D). Centrifugation was performed at 50 000 rpm and 20 °C. Radial profiles were measured at 5- or 10 min intervals at 280 nm.



Table 1 Results of AUC sedimentation velocity experiments for peptides **1**, **1_b**, **1_c** and **1_f**^a

Peptide	[c] (μM)	Solvent ^b	$s_{20,w}$ (S) (content %) ^c	ff_0 ^d	Dominant oligomerization state (theoretical $s_{20,w}$ value) ^e
1 ^f	100	H ₂ O	0.868 \pm 0.065	1.078	Dimer/trimer (0.768/1.006)
	10		0.853 \pm 0.194	1.492	Dimer/trimer (0.768/1.006)
	100	Buffer	0.972 \pm 0.088	1.213	Trimer (1.006)
	75		0.957 \pm 0.147	1.260	Trimer (1.006)
	10		0.860 \pm 0.176	1.289	Dimer/trimer (0.768/1.006)
1_b	80 (80) ^f	H ₂ O	0.524 \pm 0.049 (87)	1.521	Dimer (0.562)
			1.214 \pm 0.100 (10) ^g		Hexamer (1.170)
	30 (25)		0.514 \pm 0.098 (92) ^h	1.695	Dimer (0.562)
	80 (20)	Buffer	0.812 \pm 0.295 (50)	1.831	Trimer/tetramer (0.737/0.893)
		2.346 \pm 0.371 (26) ⁱ		Bundles	
1_c	38 (32) ^f	H ₂ O	0.989 \pm 0.080	1.161	Trimer (0.972)
	38 (27)	Buffer	1.069 \pm 0.111	1.271	Trimer (0.972)
1_f	31	H ₂ O	0.988 \pm 0.115	1.138	Trimer (0.945)

^a 50 000 rpm, scans every 5 or 10 min, 20 °C, A (280 nm), duration of experiment \sim 24 h. ^b Buffer: 50 mM potassium phosphate pH 7.0 + 50 mM NaCl. ^c When more than one species is present in the $c(s)$ distribution, their $s_{20,w}$ and percentage are given. ^d Values of ff_0 determined for the peptides dissolved in H₂O may be biased by electrostatic effects caused by the fact that charges of peptides are not shielded. ^e Theoretical values of sedimentation coefficients in the standard conditions ($s_{20,w}$) for various oligomeric forms of studied peptides are presented in Table S5_E. ^f In brackets concentration of peptide in the AUC centerpiece observed after centrifugation of large aggregates. ^g In the $c(s)$ distribution third peak is observed, $s = 2.1$ S (3%). ^h In the $c(s)$ distribution, two more peaks are observed \sim 1.1 S (7%) $i \sim$ 2.1 S (1%) (not well separated from each other and from the main peak). ⁱ Peaks are not well separated; in the $c(s)$ distribution, two more, not well separated peaks are present, spread up to \sim 10 S (max \sim 3.5 and 6.5 S, 13% and 7%, respectively); when the range 0–15 S is analyzed, large oligomers are visible, exhibiting a continuous distribution of s .

ponds to an intermediate mass between the mass of the dimer and trimer in water, and it increases slightly as the concentration of the peptide increases, which indicates that peptide **1** exists as a trimer in rapid equilibrium with smaller oligomeric forms. The value of the sedimentation coefficient obtained for 100 μM solution in AUC buffer corresponds clearly to that calculated for a trimer. The best fit to the radial concentration gradients is best described by dimer-tetramer and monomer-dimer-tetramer self-association models, but self-association models with the trimeric form, namely, monomer-trimer and monomer-dimer-trimer, show only slightly higher χ^2 and similar residual plots (Fig. S5_2 and Table S5_3[†]). The model assuming a single species indicates that the mass between the dimer and trimer and the fit of this model has a higher χ^2 value than the self-association models listed above (Table S5_3[†]). Taken together, the results from sedimentation velocity and equilibrium methods indicate that peptide **1** contains a mixture of different oligomeric forms in rapid equilibrium, but at a concentration of at least 100 μM , the trimeric form prevails.

The α,β -peptides are not well soluble in water and are even less soluble in AUC buffer. They are also much less stable in solution compared to peptide **1**, so only sedimentation velocity experiments were performed in their case. The $c(s)$ distributions obtained for water solutions of peptides **1_c** and **1_f** (Fig. 4C) contain a single peak, similar to peptide **1**. Both peptides, however, sediment faster ($s \sim$ 1 S) than the model peptide ($s \sim$ 0.9 S). These values of the sedimentation coefficient correspond to the trimeric forms of **1_c** and **1_f** (Table 1). Unlike the peptides described above, peptide **1_b** dissolved in water and in AUC buffer sediments, giving a $c(s)$ distribution with several peaks (Fig. 4C and D). Moreover, its hydrodynamic parameters are different in water and in the

AUC buffer. The species with a sedimentation coefficient of 0.5 S (attributed to dimer) is dominant in water (Fig. 4C and Table 1). Small amounts of larger oligomers are detected at 1.2 S (hexamer, 7–10%) and 2.1 S (bundles, 1–3%). The content of the oligomeric forms depends on peptide concentration in the sample, which indicates that major small species (dimer) are in equilibrium with higher oligomers.³⁷ The AUC buffer promotes the formation of large aggregates of **1_b**. Only 20% of the peptide is present as small oligomers, which can be detected at a rotor speed of 50 000 rpm (Table 1). Among them, half (50%) are in arrangements with a sedimentation coefficient of 0.85 S (trimer). Larger oligomers at 2.4 S (26%), 3.5 S (13%) and 6.5 S (7%) are also visible. The peaks observed in the $c(s)$ distribution are broad and not well separated (Fig. 4D), indicating that several types of large oligomers are in rapid equilibrium. The high value of the friction ratio, ff_0 , is another difference between **1_b** and other peptides ($ff_0 \sim$ 1.7, compared with $ff_0 \sim$ 1.25 for **1**, **1_c** and **1_f**; Table 1), which indicates deviation from the globular shape for **1_b**.

Sedimentation velocity experiments proved the presence of higher oligomers in the case of all studied peptides. The analysis of radial profiles served to determine sedimentation coefficients, which match for the trimer model **1** and for the designed peptides **1_c** and **1_f**, while peptide **1_b** showed a complicated mixture of oligomers. Therefore, $\alpha \rightarrow$ cyclic β residue replacement of outer positions c and f of the coiled-coil does not interfere significantly with its oligomerization state in solution, while the substitution of position b increases the aggregation propensity, and larger aggregates are observed in solution.

In addition to AUC experiments, the thioflavin T (ThT) assay was applied for the analysis of aggregate formation in solution.^{38–40} The ThT assay performed for model (**1**) and



α,β -peptides (**1_b**, **1_c** and **1_f**) showed increased fluorescence intensity compared to the ThT dye alone immediately after peptide dissolution (Table S6[†]), suggesting the formation of aggregates in each case. Slight enhancement of the fluorescence was observed for peptide **1** (relative fluorescence of 20.5), while α,β -peptides showed a significant increase in relative fluorescence (values of 5831, 269 and 119 for **1_b**, **1_c** and **1_f**, respectively). Subsequently, the kinetic studies were performed as the ThT fluorescence assay was often used for kinetic studies of amyloid or collagen fibrils formation.^{41–44} In case of peptide **1_b** and **1_c** sigmoidal shape of fluorescence signal was observed (Table S6[†]) with lag time of 4 h and 2 h, respectively. The lag phase indicates the unfavorable energetic state of peptide which corresponds to not sufficient amount of nuclei for the next phase. Peptides **1_b** and **1_c** showed similar kinetics of fibril formation with τ equal to 75 min and 90 min. In case of peptide **1_f** slow increase of fluorescence signal was observed, indicating that fibril formation in solution is very slow. The difference between peptides **1_b/1_c** and **1_f** can be attributed to the fact that β -residues in peptides **1_b** and **1_c** are located at the same turn of the helix, while in case of peptide **1_f**, beta-residue is on the subsequent turn. Therefore, inter-helical interactions involving β -residues are similar for **1_b** and **1_c** and different for **1_f**. Even though the peptide **1** showed a slight enhancement in the ThT emission, the fibrils were not observed under AFM and TEM. Binding to peptide **1** may be caused by unspecificity of the ThT dye, which can

bind to many structures, such as: nucleic acid, keratin, collagen.

The secondary structures of peptides **1**, **1_b**, **1_c** and **1_f** were studied by μ IR, ATR-FTIR and FT-Raman techniques both in powder form and in solution (Table S7_1[†]). The amide I band (1700–1600 cm^{-1}) characteristic of C=O stretching vibrations of the peptide backbone is sensitive to conformational changes.^{45,46} All the studied peptides possess the maximum amide I band in the range 1660–1650 cm^{-1} (Tables S7_1 and S7_2[†]) measured for powder form, which is typical for α -helical peptides. However, ATR-FTIR (Fig. S7_3[†]) and FT-Raman (Fig. S7_3[†]) spectra indicate slightly different amide I band positions for particular peptides, most likely due to the differences in the helical packing of amino acid chains.^{40,47,48} Therefore, α,β -peptides show maxima of the amide I band at lower frequencies than model peptide **1**, and the presence of *trans*-ACPC in sequence caused significant broadening of amide I bands and an increase in full width at half maximum (FWHM). More accurate information about the structure is usually provided by the second derivative and decomposition of the amide I band into sub-bands. The deconvoluted spectra were fitted with Voigtian or Lorentzian bands for ATR-FTIR and FT-Raman, respectively.⁴⁹ In the ATR-FTIR spectra of peptides **1_b**, **1_c** and **1_f**, instead of one dominant helical band in the amide I range, two components are clearly visible (Fig. 5), signifying a more complex structure. The lower band with a maximum near 1630 cm^{-1} is usually associated with

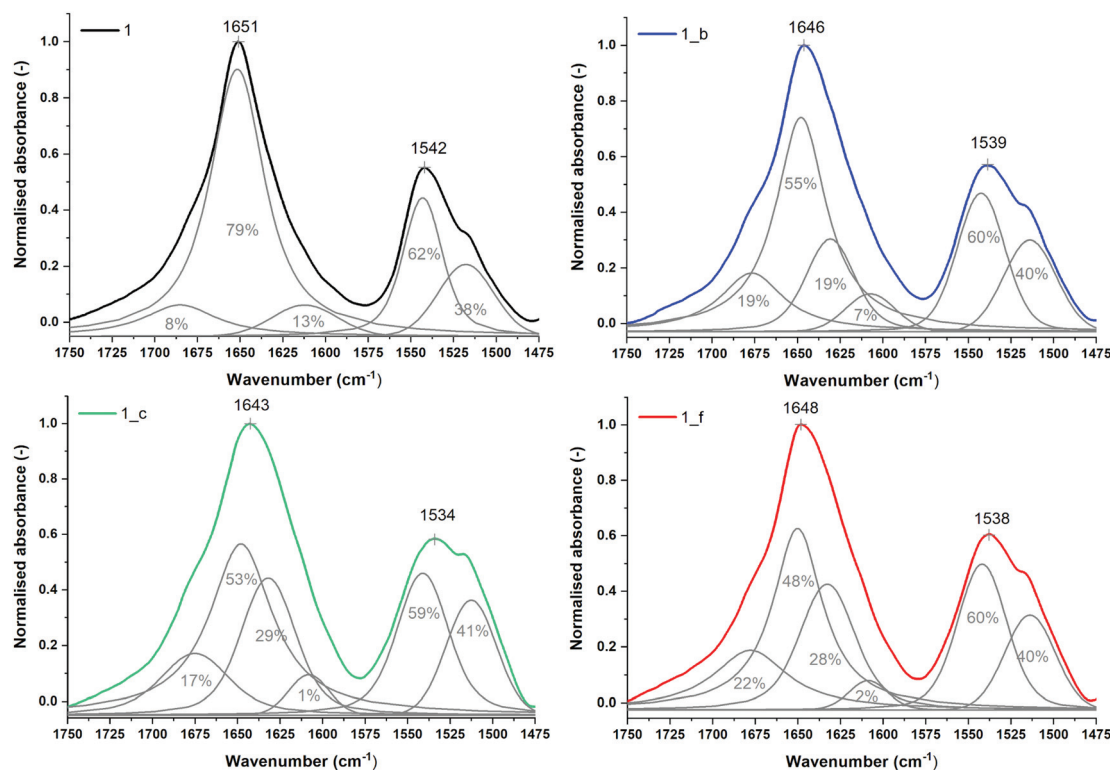


Fig. 5 Normalized ATR-FTIR spectra of powder peptides with sub-bands and percentage area values obtained from curve fitting based on their second derivatives (amide I and II regions).



aggregates.⁵⁰ We propose to assign the amide I band component at 1630 cm^{-1} to hydrogen-bonded carbonyl groups and the higher band (at 1675 cm^{-1}) to free C=O groups of exposed amino acids into the external environment. *Trans*-ACPC alone possesses an absorption band at 1691 cm^{-1} (Fig. S7_4†) and affects the area of the band with a maximum at 1675 cm^{-1} present in α,β -peptides. Deconvolution of the amide I band in the FT-Raman spectra also reveals the presence of few components in the range of $1700\text{--}1600\text{ cm}^{-1}$ (Fig. 6) with origins similar to those in the FTIR spectra. The broad peak at 1615 cm^{-1} arises from the modes of the aromatic amino acids.⁵¹

The amide II band observed in the range of $1600\text{--}1500\text{ cm}^{-1}$, derived mainly from in-plane N-H bending, also provides valuable structural information regarding secondary structures.⁵² The location of its maximum near 1542 cm^{-1} for all studied peptides is attributed to the helix structure.⁴⁰ There was no significant difference in the distribution of amide II components for the studied peptides; however, second derivatives of amide II revealed the presence of spectral features of *trans*-ACPC in the ATR-FTIR spectra of α,β -peptides (Fig. S7_3†). The amide II band is absent in the Raman spectra. In this wavenumber range, only an intense band at 1556 cm^{-1} derived from tryptophan residues is observed (Fig. 6).⁵³

No significant changes were observed in the positions of the amide bands in the infrared spectra (Fig. S7_7 and Tables

S7_1, S7_8†) obtained directly after the dissolution of the studied peptides in water, indicating preservation of the helical structure in solution. However, the shift of the maximum amide I band in the ATR-FTIR spectrum, especially evident in the case of peptide **1_b**, may suggest self-association (Table S7_8†). The ATR mode can accelerate the aggregation process due to the contact of the peptide molecules with diamond, which is why in μIR spectra, the maximum amide I mode is observed at slightly different positions. Temperature dependence spectra (Fig. S7_9†) indicate that from all studied sequences, peptide **1_b** formed the most stable structures. In the case of peptide **1_f**, the shift of the amide I maximum to higher wavenumbers suggests that it creates less unpacked structures.

No visible change in the maxima of amide I peak positions in ATR-FTIR spectra can be observed after 7 days of incubation of the peptide solutions at $37\text{ }^\circ\text{C}$, which demonstrates no alteration in the peptide secondary structures. On the other hand, in the infrared spectra of peptides **1_c** and **1_f**, the peak area of the sub-band was strongly increased at approximately 1630 cm^{-1} , which may indicate aggregate formation (Fig. S7_10 and Table S7_11†).⁵⁰ Moreover, the blueshift in the amide A maximum position ($\sim 3290\text{ cm}^{-1}$) of approximately 10 cm^{-1} suggests an increase in the strength of hydrogen interactions.^{54,55} The aggregate formation was confirmed with infrared techniques since the analysis performed after 30 days of incubation showed that for α,β -peptides, the maximum

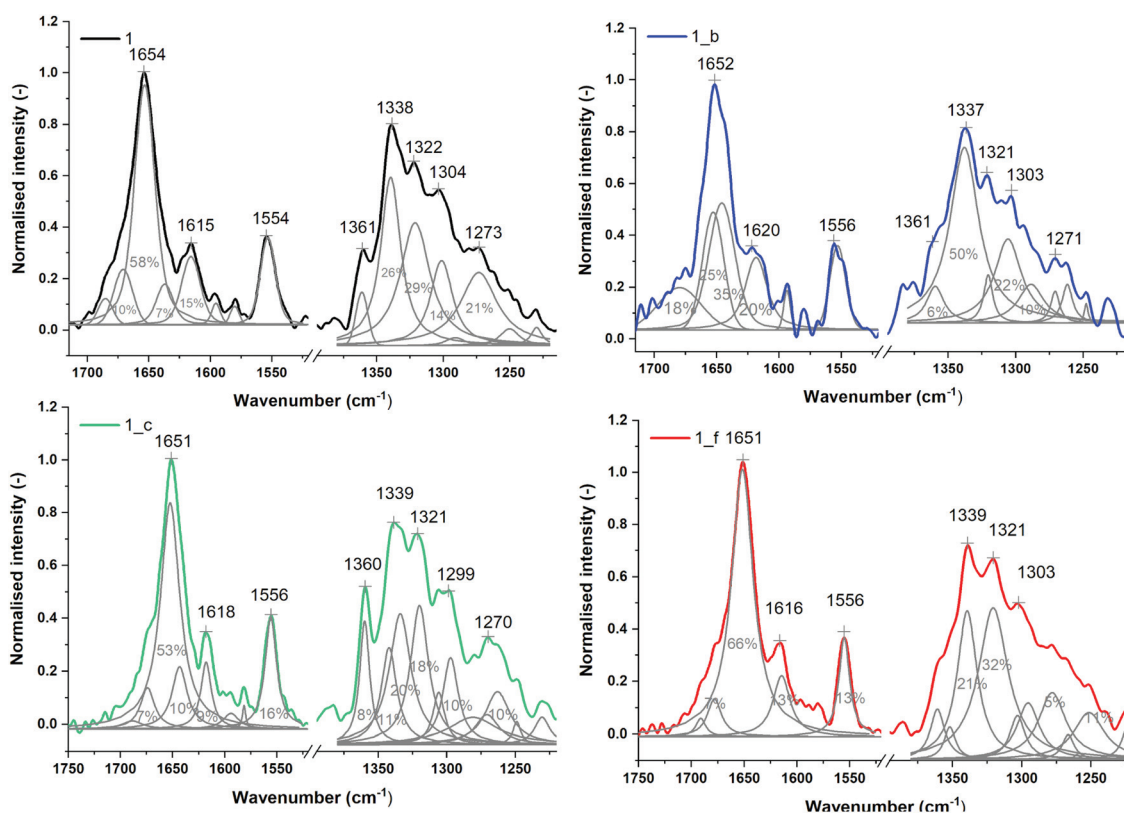


Fig. 6 Normalized FT-Raman spectra of powder peptides with sub-bands and percentage area values obtained from curve fitting based on their second derivatives (amide I and III regions).



amide I was moved from 1635 cm^{-1} to 1627 cm^{-1} , which is characteristic of long, rigid fibrils.^{50,56,57} The percentage of this sub-band was increased for samples **1**, **1_b** and **1_c** (Table S7_12†). Only peptide **1_f** did not show a significant decline in percentage (2.43%).

All peptides demonstrated spectral features characteristic of helical secondary structures, both in powder and in water solutions. Peptides **1_b**, **1_c** and **1_f** containing cyclic β -amino acids tend to adopt more compact structures than model peptide **1**. Additionally, changes in the band positions of α,β -peptides after several days of incubation may indicate self-association and aggregate formation. Similar shifting behaviors of amide I position and self-aggregation were observed for phenol-soluble modulins $\alpha 3$ (PSM $\alpha 3$) from *Staphylococcus aureus*⁵⁸ and *de novo* designed peptides⁵⁹ forming a cross- α architecture.

Nanostructure characterization

We used atomic force microscopy (AFM) and transmission electron microscopy (TEM) to observe the results of peptide self-aggregation. Basing on AFM measurements, we established that peptide **1** was not able to self-assemble molecularly

on a mica surface (Fig. S8_1†). The aggregation process was not observed even after two weeks of sample incubation. On the other hand, α,β -peptides **1_b**, **1_c** and **1_f** self-assemble immediately, and over a period of time, the formation of long fibrils can be observed (Fig. 7). After one hour of adsorption on the mica surface, we detected the formation of globular structures, which created fibrils after a few hours. After several days of incubation, the fibrils became crosslinked, and the average diameter of the structures changed from 3 to 6.4 nm. We presume that such a phenomenon may be caused by secondary nucleation of the aggregation process, similar to the one observed in amyloid proteins.⁶⁰ The self-association process occurs faster for **1_b** than for **1_c** and **1_f**; however, in all cases, fibrils are formed even at low concentrations ($<80\ \mu\text{M}$). Morphological differences between the formed fibers, such as the smoothness of the fiber edges, can also be observed (Fig. 7). The average heights of the helical structures for all peptides ranged from 1.35 to 4.34 nm (Fig. S8_2†). The tallest structures were formed by peptide **1_c**, directly after adsorption on the mica surface (Fig. 7 and S8_2†). As the incubation time increased, the height of the peptides decreased, with one exception: the height of peptide **1_f** increased from

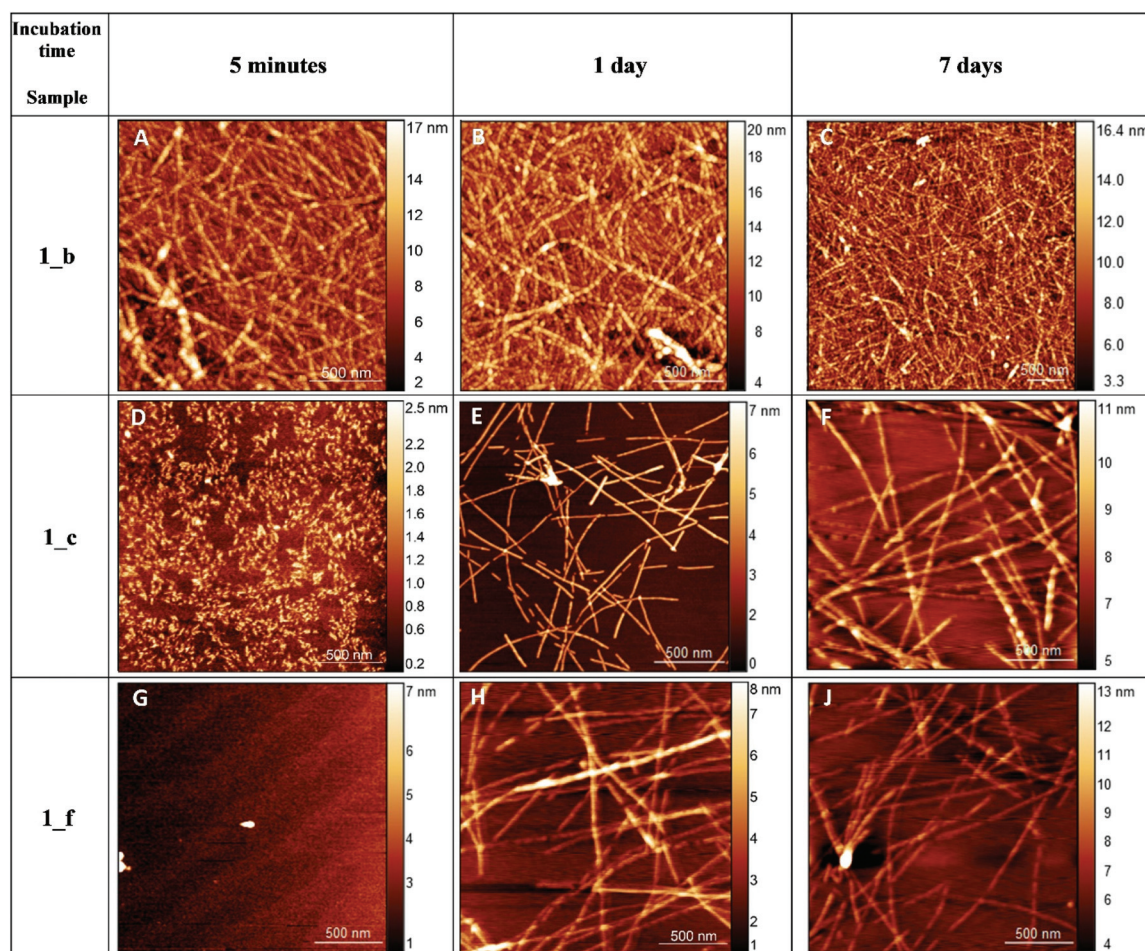


Fig. 7 AFM images of peptides **1_b**, **1_c** and **1_f** on the mica surface after various incubation times ($C_{\text{peptide}} = 1.3\text{ mM}$).



1.35 to 1.53 nm. The observed fluctuation in the height of fibrils can be a sign of different stages of peptide aggregation, as mentioned above.⁶¹

TEM analysis confirmed the pattern of nanostructure formation in water, starting from globular aggregates to fibrils, after a short period of time (Fig. 8). TEM allows us to determine more accurate values of average diameter, which are 10.42 ± 2.72 nm for **1_b**, 10.18 ± 1.93 nm for **1_c** and 12.08 ± 2.65 nm for **1_f** (Fig. S9†). The average contour length of fibrils amounts to 412 ± 61.53 nm for **1_c** and 49.8 ± 27.9 nm for **1_f** (Fig. S9†). Fibrils formed by peptide **1_b** were too long and tangled to determine their length. In terms of fibril architecture, nanostructures formed by peptide **1_f** differ from structures assembled by peptides **1_b** and **1_c** (Fig. 8); in the case of peptide **1_f**, there are visible small, globular aggregates that interact with each other to form fibril-like structures. However,

after 7 days of incubation, long fibrils can be also visible for peptide **1_f** (Fig. S10†). The fibers created by peptides **1_b** and **1_c** are more regular, showing a twisted rope-like hierarchical structure. It appears that the fibers form predepositions as several of the fibers lay across the top of each other.

AFM and TEM analyses of the studied α,β -peptides give consistent results in terms of the type, rate and amount of nanostructure formation. It can be concluded that even at a low concentration, globular aggregates form, and after a short period of time form long fibrils. The dissimilarities in nanostructure morphology, especially for peptide **1_f**, may arise from the differences in sample adsorption (mica surface in AFM *versus* carbon grid in TEM).

The X-ray diffraction experiments revealed presence of ordered microstructure in the dried sample of peptide **1_f** deposited on SiN window. Diffraction pattern (Fig. 9A) shows

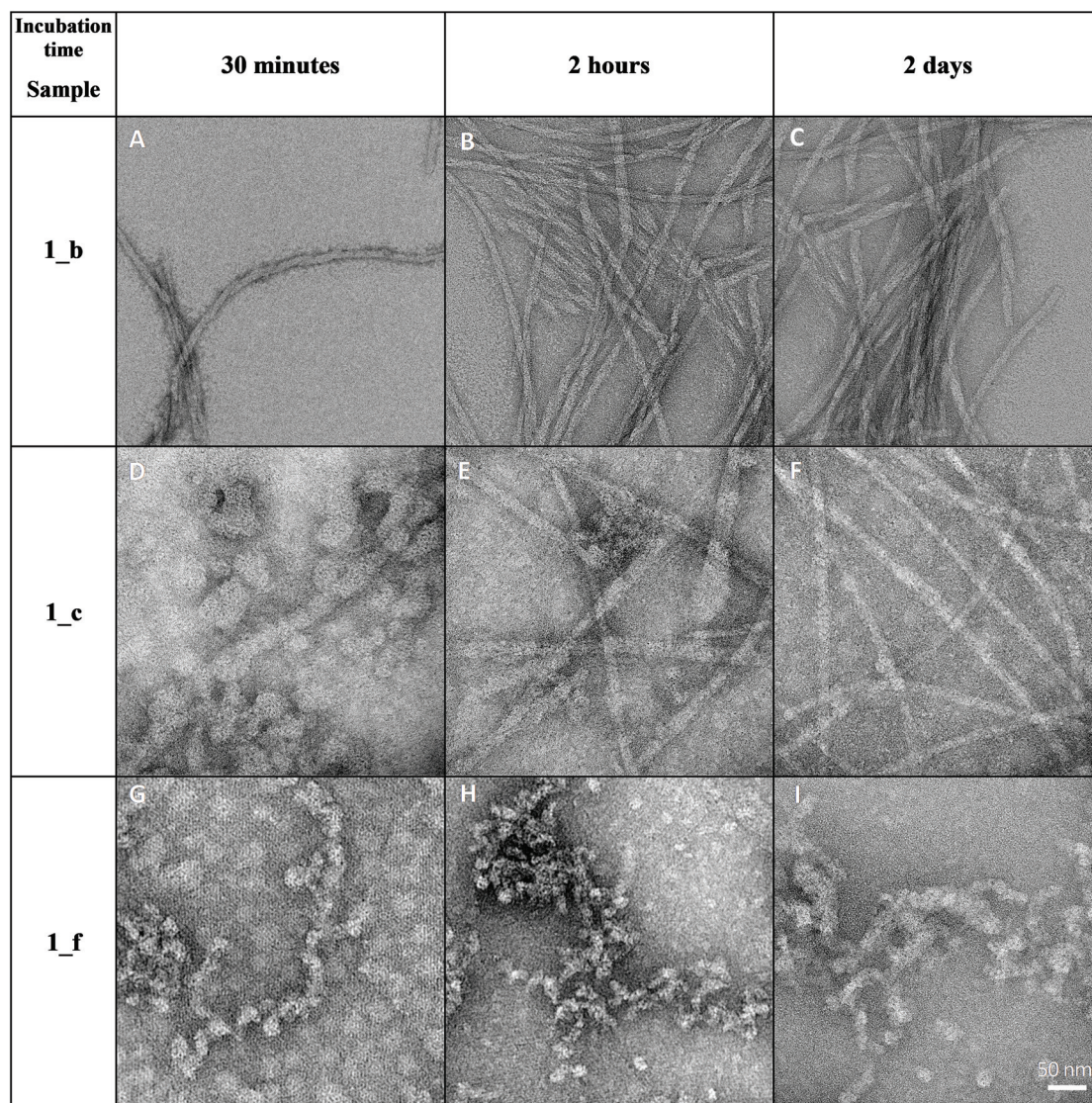


Fig. 8 TEM images of peptides **1_b**, **1_c** and **1_f** on carbon grid surface after various incubation times ($c_{\text{peptide}} = 1.3$ mM). Images registered at the magnification of 50 nm.



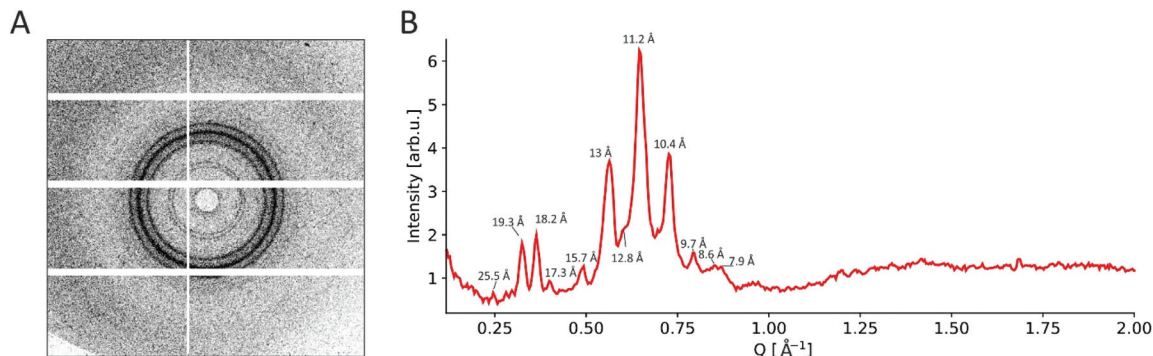


Fig. 9 X-ray diffraction pattern of the peptide **1_f** sample at 80 μM dried on SiN window. X-ray diffraction pattern of non-aligned sample (A). Peak profile of X-ray diffraction pattern with calculated d -spacing of corresponding peaks. $Q = 4\pi \sin(\theta)/\lambda$, where θ is diffraction angle and λ is the X-ray radiation wavelength (B).

multiple co-centric diffraction rings characteristic for crystallites or fibers without preferential orientation. The diffraction profile was obtained by integration of full diffraction image (Fig. 9B). It shows multiple diffraction peaks of various intensity in the Q vector range from 0.25 \AA^{-1} to 1 \AA^{-1} . In general, such a diffraction pattern is characteristic for the α -helical assemblies. The peak corresponding to d -spacing equal to 10.4 \AA is characteristic also for coiled-coil peptides assemblies heptad repeat pattern.⁶² The same value was observed by the use of the electron cryo-transmission microscopy for *de novo* designed synthetic coiled-coil peptides.⁶³ In this case, those correspond to the packing parameter of the coiled-coil dimer into the larger fiber bundle. Observed diffraction pattern of peptide **1_f** is different from the typical pattern for cross- β structure of amyloid fibrils.⁶⁴ Those fibers has characteristic diffraction rings at 10–11 \AA and 4.7 \AA d -spacings, corresponding to the spacing between β -strands and spacing between β -sheets, respectively.

Recent studies also showed that some peptides can form cross- α structure fibrils.^{58,65} In those structures α -helices aligns in the similar fashion to the β -strands of amyloid fibers. X-ray diffraction studies of the wild-type phenol-soluble modulin $\alpha 3$ (PSM $\alpha 3$) peptides shows characteristic diffraction patterns with d -spacing calculated for main reflections equal to: 10.5 \AA , 12 \AA and 25 \AA . In the diffraction pattern of peptide **1_f** fibrils a strong peaks at 25 \AA is absent. There is only very weak diffraction peak at 25.5 \AA . This phenomena suggests that peptide **1_f** could form some coiled-coil structures than cross- α or cross- β structures, however further detailed studies are needed to resolve three-dimensional structure of the peptide **1_f** assemblies.

Conclusions

A methodology for the rational construction of helix-containing nanofibers using the modification of coiled-coil structure outer positions with helix-promoting *trans*-ACPC residue was elaborated. Incorporation of one *trans*-ACPC residue per

heptad induced the formation of helical structures only if this substitution did not interfere with further coiled-coil formation; *i.e.*, only the outer b , c and f positions were available for this modification. The formation of helices was confirmed by various techniques, including CD, NMR and FT-IR/Raman. Importantly, significantly higher conformational stability was shown for *trans*-ACPC-modified peptides in comparison to model peptide **1**. Subsequently, the formation of aggregates was confirmed in solution using AUC and ThT assays. In the cases of two modified peptides, namely, **1_c** and **1_f**, the formation of trimers was evidenced in solution, which is analogous to starting peptide **1**. Again, the propensity of **1_c** and **1_f** for trimer formation was higher than that of unmodified peptide **1**. The behavior of peptide **1_b** in solution was significantly different, as a large number of larger structures were detected. Subsequently, peptides were studied in solid state on mica and carbon surfaces. Well-formed nanofibrils were observed using both AFM and TEM techniques for peptides **1_b**, **1_c** and **1_f**. At that level of molecular organization, the difference between *trans*-ACPC-substituted and original coiled-coil peptides was the most substantial, as model peptide **1** did not form any nanostructures. The velocity of nanostructure formation in the studied peptides was different, which allowed us to observe the initial globular aggregates preceding the creation of nanofibers in some cases. Therefore, it could be assumed that the pathway of fibril formation includes the following stages: helices \rightarrow oligomers \rightarrow bundles \rightarrow globular aggregates \rightarrow nanofibrils. Importantly, individual features of the studied peptide sequences could be attributed to the achievement of a particular level of organization, which makes the design process rational. Structural constraints introduced by rigid *trans*-ACPC residues contribute to a higher propensity of the peptides to form helix structures. The residues in positions a/d and e/g in the heptad are responsible for coiled-coil formation (trimers). The formation of larger aggregates is driven by the introduced hydrophobic surface delivered by the cyclopentane side chains of *trans*-ACPC residues. We propose the formation of a zipper-like interaction between helices by analogy to the leucine zippers observed in transcriptional regu-



lators.⁶⁶ We consider that the presented methodology will constitute a significant step towards a better understanding of beta-amino acid-containing peptide properties and will provide a basis for future discoveries of higher-order structures of peptides from this class.

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*S*)-2-aminocyclopentanecarboxylic acid (*trans*-ACPC) was purchased from Synnovator, Inc. Described peptides were obtained with an automated solid-phase peptide synthesizer (Biotage® Initiator + Alastra™) using rink amide AM resin (loading: 0.59 mmol g⁻¹). Fmoc deprotection was performed using 20% piperidine in DMF for 3 + 10 min. A double-coupling procedure was achieved with 0.5 M solution of DIC and 0.5 M solution of OXYMA (1 : 1) in DMF, for α -amino acids 2 \times 15 min and for β -amino acids 30 min, at 75 °C. Acetylation reaction was carried out using NMP/DIPEA/acetic anhydride (80 : 15 : 5) mixture. Cleavage of the peptides from the resin was accomplished with the mixture of TFA/TIS/H₂O (95 : 2.5 : 2.5) after 3 h of shaking. The crude peptide was precipitated with ice-cold Et₂O and centrifuged (10 000 rpm, 15 min, 4 °C). Peptides were purified using preparative HPLC (Knauer Prep) with a C18 column (Thermo Scientific, Hypersil Gold 12 μ , 250 mm \times 20 mm) with water/acetonitrile (0.05% TFA) eluent system.

Analytical high-performance liquid chromatography (HPLC)

Analytical HPLC was performed using Kinetex 5 μ EVO C18 100A 150 \times 4.6 mm column. Program (eluent A: 0.05% TFA in H₂O, eluent B: 0.05% TFA in acetonitrile, flow 0.5 mL min⁻¹): A: $t = 0$ min, 90% A; $t = 30$ min, 10% A.

Mass spectrometry (MS)

Peptides were studied by WATERS LCT Premier XE System consisting of high resolution mass spectrometer with a time of flight (TOF).

Analytical ultracentrifugation (AUC)

Analytical ultracentrifugation was run at 20 °C on Beckman Optima XL-I with 4- or 8-position An-Ti rotor and UV detection at 280 nm, in a double-sector 1.2 cm cells with charcoal-filled epon centrepieces and sapphire windows. Lyophilized peptides were dissolved in water or in AUC buffer (50 mM phosphate buffer pH 7.0, 100 mM NaCl) directly before measurements. Peptides concentration (10–140 μ M) was determined spectrophotometrically (Jasco V-650 spectrophotometer) from absorption values at 280 nm using the extinction coefficients $\epsilon = 6990$ M⁻¹ cm⁻¹ calculated from the amino acid composition by ProtParam.⁶⁷ Sedimentation velocity experiments were performed at 50 000 rpm. Radial absorption scans of peptide concentration profiles were measured at 5 or 10 min intervals.

390 μ l of a peptide sample and 400 μ l of a reference solution (water or AUC buffer) were loaded into the right and the left sector of the cell. The sedimentation velocity data were analysed using program SEDFIT with a continuous sedimentation coefficient distribution model, $c(s)$, based on the Lamm equation.^{68,69} Integration of the $c(s)$ peaks provided the signal weighted average sedimentation coefficients (s) and corresponding standard sedimentation coefficients $s_{20,w}$ (referring to the water and 20 °C). The density and viscosity of the AUC buffer were calculated using Sednterp program.⁷⁰ The partial specific volumes of peptides were calculated from the partial specific volumes obtained in the Sednterp program (based on amino acid composition excluding ACPC residues, acetyl group at the N-terminus and NH₂ group at the C-terminus) and from partial specific volumes of the ACPC residues ($\bar{v} = 0.929207$ cm³ g⁻¹) and of the end groups (Ac and NH₂, $\bar{v} = 0.813588$ cm³ g⁻¹) obtained according to the method of Durchschlag and Zipper⁷¹ (Table S5_D†). Multispeed sedimentation equilibrium experiments were performed at 26 000, 32 000, and 50 000 rpm. 150 μ l of a peptide samples and 160 μ l of the AUC buffer (as a reference) were placed in the cells. The radial concentration gradients were collected every 3 h at the radial intervals of 0.001 cm, until the sedimentation equilibrium was attained (25–35 h per each speed). The data were analysed by means of nonlinear regression using SEDPHAT program, according to single species model, monomer- n -mer association or monomer- n -mer- m -mer association, with a global fitting to all equilibrium profiles obtained at all three rotor speeds.⁷² Theoretical values of sedimentation coefficients in the standard conditions ($s_{20,w}$) for various oligomeric forms of studied peptides were calculated using the formula $s_n = s_1 \times n^{2/3}$, where s_1 is sedimentation coefficient for a monomer.⁷³ First, $s_{20,w}$ values for trimeric form ($n = 3$) were obtained in the SEDFIT program using its' $M(s)$ calculator, and introducing constraints that molecular mass calculated for the particular $s_{20,w}$ value must be equal to the mass of the peptide calculated from its amino acid sequence. Values f/f_0 that are necessary for these calculations were $f/f_0 = 1.237$, 1.676 and 1.271 for peptides **1**, **1_b**, and **1_c** and **1_f**, respectively, hence taken from the Table 1. In the next step $s_{20,w}$ for other oligomeric forms were calculated using the formula $s_n = s_1 \times n^{2/3}$.

Circular dichroism (CD)

CD spectra were recorded on JASCO J-815 at 20 °C between 260 and 190 nm in water with following parameters: 0.2 nm resolution, 1.0 nm bandwidth, 20 mdeg sensitivity, 0.25 s response, 50 nm min⁻¹ scanning speed, 5 scans, 0.02 cm cuvette path length. The CD spectra of the solvents alone were recorded and subtracted from the raw data. For peptides dissolved in water pH was adjusted to 7. Typically, the samples were prepared by dilution of peptides stock solution to obtain peptide concentration of 80 μ M. The CD intensity is given as mean residue molar ellipticity ($[\theta]$ [deg cm² dmol⁻¹]). The melting temperatures of studied peptides were determined by method described previously.⁷⁴



Temperature denaturation measurements using CD

To examine the thermal unfolding of the peptides, stock solutions were diluted to 80 μM and three types of measurements were performed: (1) in the scope 190–240 nm; (2) at 208 nm in water and (3) at 208 nm with 3 M GuHCl. The temperature was increased from 4 to 98 $^{\circ}\text{C}$ in increments of 2 $^{\circ}\text{C}$. Ellipticity measurements were recorded with 1 mm path length cuvette, others parameters remained unchanged.

Nuclear magnetic resonance (NMR)

The NMR experiments were performed on Bruker AvanceTM spectrometer operating at 600.58 MHz for ^1H , equipped with a 5.0 mm PA BBO probe. The NMR spectra at temperature 298 K were recorded in CD_3OH . 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 solvent OH signal using 3-9-19 pulse sequence with gradients. Typical TOCSY – homonuclear Hartman (Hahn transfer using 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 TOCSY and 80 scans for 2D NOESY. Mixing times were set at 80 ms and 200 ms for TOCSY and 2D NOESY experiments, respectively. Processing and analysis of the NMR data were performed on Topspin 3.2 (Bruker BioSpin) software.

NMR structure determination

NMR structure determination was performed using Xplor-NIH v. 2.41 program.⁷⁵ NMR derived inter-proton contacts were classified by standard method with upper distance limits: strong 2.5 \AA (s), medium 3.5 \AA (m) and weak 5 \AA (w), and the lower distance limit was set to 1.8 \AA . Initially, 50 random conformations of a peptide were generated. Restraints derived from NOE contacts and ^3J coupling constants of amide protons were applied. Standard protocol for NMR structure calculations implemented in Xplor-NIH was used with the following steps: (a) high temperature dynamics (3500 K, 800 ps or 8000 steps), (b) 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); (c) gradient minimization of final structure. Finally, top lower energy structures were superimposed.

Vibrational spectroscopy

Attenuated total reflectance – Fourier-transform infrared (ATR-FTIR) studies were performed on FT-IR spectrometer (Thermo Scientific, USA) with Golden Gate Mk II ATR Accessory with Heated Diamond Top-plate (PIKE Technologies). ATR-FTIR spectra were obtained in the range of 3600–150 cm^{-1} . For each spectrum 128 interferograms was coadded with 4 cm^{-1} resolution. Directly before sampling, the background spectrum of diamond/air was recorded as a refer-

ence (512 scans, 4 cm^{-1}). The spectra of 1.3 mM peptides solutions were registered in the temperature range 25–95 $^{\circ}\text{C}$ (77–203 $^{\circ}\text{F}$) with a 2 $^{\circ}\text{C}$ step. Additionally, air-dried films were obtained. Initially, 10 μl of peptide aqueous solution was dropped directly on the diamond surface and was allowed to dry out. At the time, spectra were recorded at 5 minutes intervals. Infrared (IR) spectra were collected using integrated infrared microscope FTIR iN10 (Thermo Scientific, USA) equipped with a liquid nitrogen cooled mercury cadmium telluride (MCT-A) detector. All spectra were acquired in the transmission mode (10 \times 10 image size, 50 micron aperture, 50 micron step size, 64 scans per point, resolution of 4 cm^{-1}). A drop of 10 μl of each sample was deposited onto BaF_2 windows and air dried prior to imaging. Measurements were conducted at room temperature, approximately 22 $^{\circ}\text{C}$ (72 $^{\circ}\text{F}$). Raman spectra were measured using NXR FT-Raman spectrometer with MicroStage extension equipped with Nd:YVO₄ laser (1064 nm, 500 mW) as an excitation source and InGaAs detector. FT-Raman were recorded over a range of 3700–150 cm^{-1} at 4 cm^{-1} resolution. In each case, an average of 1024 scans were collected. Only samples in powder form were tested. All spectra were analyzed using the OriginPro 9.0 (OriginLab Corporation, USA). The analysis included spectra baseline correction, smoothing using the Savitzky–Golay filter (parameters dependent on the type of spectra), normalization of spectra relative to selected bands and deconvolution into subcomponents using the Lorentz function based on second derivative spectra. ATR-FTIR spectra were initially preprocessed using OMNIC 8.3.103.

Atomic force microscopy (AFM)

AFM images were collected using a MultiMode atomic force microscope with a NanoScope IV controller (Digital Instruments, Santa Barbara, CA). The topography was measured in tappingTM mode (TM). The tip, a force modulation etched silicon probe (Veeco Metrology, Sunnyvale, CA), had a constant of 23 N m^{-1} , a resonance frequency of 296–351 kHz, a nominal tip radius of curvature 5 nm and a cantilever length of 110–140 μm . NCLR and TESP-SS cantilevers were used. 10 μl of the 1.3 mM peptide solution in water was applied to freshly cleaved mica sheet and after 1 minute of adsorption the sheet was rinsed with deionized water. Time of measurements: directly after one minute of adsorption on the mica surface (about 5 minutes), 1 day and 7 days. Samples were incubated at 25 $^{\circ}\text{C}$. The resolution of the AFM scans was 512 \times 512 pixels, which produces topographic images of the studied peptide on the mica surface (2 $\mu\text{m} \times$ 2 μm). Images were collected several times across the mica surface. The images were analyzed and processed using Gwyddion software (Czech Metrology Institute). We used first-order flattening tool – Gaussian and sharpening filters on the height images. The surface images (3D AFM) were composed of small parts of each image. The average height of nanostructures was established by performing statistical analysis of the AFM images. To determine the distribution of acquired data two normality tests Shapiro–Wilk and Kolmogorov–Smirnov were performed.



All data describing vertical distance of peptides come from normal distribution (p -value <0.05). Artifacts, like curvature distortion (blowing effects), were considered during morphological analysis.

Transmission electron microscopy (TEM)

The samples were suspended in water to the concentration of 1.3 mM and incubated at 37 °C. After 30 minutes, 2 and 48 hours from the incubated samples a small amount of suspension was taken and spotted on a carbon on copper grid. After 1 minute, an excess of the material was removed and 2% uranyl acetate was applied to the grid to stain negative peptide structures. After another minute, an excess of the staining was collected and the grid was allowed to dry under atmospheric conditions. Imaging was performed using transmission electron microscope Tecnai TF 20 X-TWIN (200 kV).

Thioflavin T (ThT) fluorescence assay

ThT powder was dissolved in MilliQ® to final concentration 10 μM. ThT solution was dissolved in 50 mM phosphate buffer (pH = 7) to final concentration 50 mM. The 90 μL of ThT buffer was mixed with 10 μL of peptide solution (concentration 1.3 mM) in the 96 wells plate. Samples were measured on the SpectraMax® Gemini™ XPS Microplate (Molecular Devices LLC). The plate reader was heated to 37 °C. The excitation wavelength was set at 440 nm and the emission was recorded in the range from 450 to 600 nm (slit widths 3 nm). The examined peptides were measured directly after dissolving. The measurements were repeated 3 times.

Kinetic measurements were done for the samples containing 10 μL of 1 mg mL⁻¹ peptide and 90 μL of 50 μM ThT solution mixed in a 96-well plate. The fluorescence of each well was read by microplate reader CLARIOstar Plus, BMG LABTECH at 25 °C with 30 s shaking, every 30 min, during 24 hours measurements. The excitation wavelength was set at 440 nm and emission at 480 nm. Each group of experiment contained six parallel samples and the data were averaged after measurements.

The aggregation kinetic curves obtained for studied peptides represent classic example of a ThT-monitored kinetic curves and so were fitted by the sigmoidal curves using the following eqn (1):

$$Y = y_i + m_i t + \frac{y_f + m_f t}{1 + e^{-[(t-t_{1/2})/\tau]}} \quad (1)$$

where, Y is the fluorescence intensity as a function of time t , y_i and y_f are the intercept of the initial and final baselines with the y -axis, m_i and m_f are slopes of the initial and final baselines, $t_{1/2}$ is the time needed to reach halfway through the elongation phase and τ is the elongation time constant. The apparent rate constant, k_{app} , for the growth of fibrils is given by $1/\tau$, and the lag time is defined as $t_{lag} = t_{1/2} - 2\tau$

Wide-angle X-ray scattering (WAXS)

The fiber diffraction experiments were performed on laboratory SAXS/WAXS Xeuss 2.0 system (Xenocs, Sassenage, France)

equipped with MetalJet D2 microfocuss X-ray source ($\lambda = 0.134$ nm) with a liquid gallium alloy target (Excillum AB, Kista, Sweden). 1 mL of fibrillated peptide 1_f sample at concentration of 80 μM in H₂O were centrifuged for 10 minutes at 16 000g. The supernatant was discarded and the pellet was resuspended in 30 μL of H₂O. About 20 μL of obtained sample was placed dried in 2 μL portions on SiN windows (Xenocs, Sassenage, France). The deposited on SiN sample was then placed on sample holder and measured in vacuum for 10 minutes. Sample-to-detector distance, as well as diffraction range, were calibrated using diffraction pattern of silver behenate as a reference sample.⁷⁶ Obtained X-ray diffraction images were processed using Foxtrot package⁷⁷ and background from empty SiN window was also subtracted. The real space d -spacing values were calculated from the formula: $d = 2\pi/s$, where s is a position of the peak maximum in reciprocal (Q) space.

Abbreviations

trans-ACPC *trans* (1*S*,2*S*)-2-Aminocyclopentanecarboxylic acid
ThT Thioflavin T

Funding sources

The work was financially supported by the National Science Centre, Poland, Grant No. 2017/26/D/ST5/00341 (to M. S.)

Conflicts of interest

Authors declare no potential conflict of interest.

Acknowledgements

The authors would like to gratefully acknowledge National Science Centre, Poland for their financial support through Grant No. 2017/26/D/ST5/00341. M. S. would like to thank to Prof. Małgorzata Kotulska for fruitful discussions. This work was partially performed in the NanoFun laboratories co-financed by the European Regional Development Fund within the Innovation Economy Operational Program, the Project No. POIG.02.02.00-00-025/09.

References

- 1 E. De Santis and M. G. Ryadnov, Peptide Self-Assembly for Nanomaterials: The Old New Kid on the Block, *Chem. Soc. Rev.*, 2015, **44**, 8288–8300.
- 2 E. H. C. Bromley and K. J. Channon, Alpha-Helical Peptide Assemblies: Giving New Function to Designed Structures, *Prog. Mol. Biol. Transl. Sci.*, 2011, **103**, 231–275.



- 3 L. E. R. O'Leary, J. A. Fallas, E. L. Bakota, M. K. Kang and J. D. Hartgerink, Multi-Hierarchical Self-Assembly of a Collagen Mimetic Peptide from Triple Helix to Nanofibre and Hydrogel, *Nat. Chem.*, 2011, **3**, 821–828.
- 4 D. M. Raymond and B. L. Nilsson, Multicomponent Peptide Assemblies, *Chem. Soc. Rev.*, 2018, **47**, 3659–3720.
- 5 P. B. Harbury, T. Zhang, P. S. Kim and T. Alber, A Switch between 2-Stranded, 3-Stranded and 4-Stranded Coiled Coils in Gcn4 Leucine-Zipper Mutants, *Science*, 1993, **262**, 1401–1407.
- 6 J. M. Mason and K. M. Arndt, Coiled Coil Domains: Stability, Specificity, and Biological Implications, *ChemBioChem*, 2004, **5**, 170–176.
- 7 Y. B. Yu, Coiled-Coils: Stability, Specificity, and Drug Delivery Potential, *Adv. Drug Delivery Rev.*, 2002, **54**, 1113–1129.
- 8 P. S. P. Wang and A. Schepartz, β -Peptide Bundles: Design. Build. Analyze. Biosynthesize, *Chem. Commun.*, 2016, **52**, 7420–7432.
- 9 D. N. Woolfson, Coiled-Coil Design: Updated and Upgraded, *Subcell. Biochem.*, 2017, **82**, 35–61.
- 10 C. Xu, R. Liu, A. K. Mehta, R. C. Guerrero-Ferreira, E. R. Wright, S. Dunin-Horkawicz, K. Morris, L. C. Serpell, X. Zuo, J. S. Wall and V. P. Conticello, Rational Design of Helical Nanotubes from Self-Assembly of Coiled-Coil Lock Washers, *J. Am. Chem. Soc.*, 2013, **135**, 15565–15578.
- 11 J. M. Fletcher, R. L. Harniman, F. R. Barnes, A. L. Boyle, A. Collins, J. Mantell, T. H. Sharp, M. Antognozzi, P. J. Booth, N. Linden, M. J. Miles, R. B. Sessions, P. Verkade and D. N. Woolfson, Self-Assembling Cages from Coiled-Coil Peptide Modules, *Science*, 2013, **340**, 595–599.
- 12 Y. Wu and J. H. Collier, α -Helical Coiled-Coil Peptide Materials for Biomedical Applications, *Wiley Interdiscip. Rev.: Nanomed. Nanobiotechnol.*, 2017, **9**, 1424.
- 13 D. Seebach, A. K. Beck and D. J. Bierbaum, The World of β - and γ -Peptides Comprised of Homologated Proteinogenic Amino Acids and Other Components, *Chem. Biodiversity*, 2004, **1**, 1111–1239.
- 14 D. E. Mortenson, D. F. Kreidler, N. C. Thomas, I. A. Guzei, S. H. Gellman and K. T. Forest, Evaluation of β -Amino Acid Replacements in Protein Loops: Effects on Conformational Stability and Structure, *ChemBioChem*, 2018, **19**, 604–612.
- 15 J. L. Price, W. S. Horne and S. H. Gellman, Structural Consequences of β -Amino Acid Preorganization in a Self-Assembling α/β -Peptide: Fundamental Studies of Foldameric Helix Bundles, *J. Am. Chem. Soc.*, 2010, **132**, 12378–12387.
- 16 R. D. Gopalan, M. P. Del Borgo, A. I. Mechler, P. Perlmutter and M.-I. Aguilar, Geometrically Precise Building Blocks: the Self-Assembly of β -Peptides, *Chem. Biol.*, 2015, **22**, 1417–1423.
- 17 K. Kulkarni, N. Habila, M. P. Del Borgo and M.-I. Aguilar, Novel Materials From the Supramolecular Self-Assembly of Short Helical β 3-Peptide Foldamers, *Front. Chem.*, 2019, **7**, 70.
- 18 W. S. Horne, J. L. Price, J. L. Keck and S. H. Gellman, Helix Bundle Quaternary Structure from α/β -Peptide Foldamers, *J. Am. Chem. Soc.*, 2007, **129**, 4178–4180.
- 19 W. S. Horne, M. D. Boersma, M. A. Windsor and S. H. Gellman, Sequence-Based Design of α/β -Peptide Foldamers That Mimic BH3 Domains, *Angew. Chem., Int. Ed.*, 2008, **47**, 2853–2856.
- 20 S. H. Yoo and H.-S. Lee, Foldectures: 3D Molecular Architectures from Self-Assembly of Peptide Foldamers, *Chem. Res.*, 2017, **50**, 832–841.
- 21 Z. Yang, G. Liang, M. Ma, Y. Gao and B. Xu, In Vitro and In Vivo Enzymatic Formation of Supramolecular Hydrogels Based on Self-Assembled Nanofibers of a β -Amino Acid Derivative, *Small*, 2007, **3**, 558–562.
- 22 J. Mangelschots, M. Bibian, J. Gardiner, L. Waddington, Y. Van Wanseele, A. Van Eeckhaut, M. M. Acevedo, B. Van Mele, A. Madder, R. Hoogenboom and S. Ballet, Mixed α/β -Peptides as a Class of Short Amphiphathic Peptide Hydrogelators with Enhanced Proteolytic Stability, *Biomacromolecules*, 2016, **17**, 437–445.
- 23 W. C. Pomerantz, V. M. Yuwono, R. Drake, J. D. Hartgerink, N. L. Abbott and S. H. Gellman, Lyotropic Liquid Crystals Formed from AHC-Rich β -Peptides, *J. Am. Chem. Soc.*, 2011, **133**, 13604–13613.
- 24 J. Nanda and A. Banerjee, A Gel-Based Trihybrid System Containing Nanofibers, Nanosheets, and Nanoparticles: Modulation of the Rheological Property and Catalysis, *Soft Matter*, 2012, **8**, 3380–3386.
- 25 L. K. A. Pils and O. Reiser, Alpha/Beta-Peptide Foldamers: State of the Art, *Amino Acids*, 2011, **41**, 709–718.
- 26 M. Szczyzyk, E. Weglarz-Tomczak, P. Fortuna, A. Krzyszton, E. Rudzinska-Szostak and L. Berlicki, Controlling the Helix Handedness of $\alpha\beta$ -Peptide Foldamers through Sequence Shifting, *Angew. Chem., Int. Ed.*, 2017, **56**, 2087–2091.
- 27 I. M. Mandity, E. Weber, T. A. Martinek, G. Olajos, G. K. Toth, E. Vass and F. Fulop, Design of Peptidic Foldamer Helices: A Stereochemical Patterning Approach, *Angew. Chem., Int. Ed.*, 2009, **48**, 2171–2175.
- 28 K. Ananda, P. G. Vasudev, A. Sengupta, K. M. P. Raja, N. Shamala and P. Balaram, Polypeptide Helices in Hybrid Peptide Sequences, *J. Am. Chem. Soc.*, 2005, **127**, 16668–16674.
- 29 M.-R. Lee, N. Raman, S. H. Gellman, D. M. Lynn and S. P. Palecek, Incorporation of β -Amino Acids Enhances the Antifungal Activity and Selectivity of the Helical Antimicrobial Peptide Aurein 1.2, *ACS Chem. Biol.*, 2017, **12**, 2975–2980.
- 30 C. M. Goodman, S. Choi, S. Shandler and W. F. DeGrado, Foldamers as Versatile Frameworks for the Design and Evolution of Function, *Nat. Chem. Biol.*, 2007, **3**, 252–262.
- 31 R. R. Araghi, C. Baldauf, U. I. M. Gerling, C. D. Cadicamo and B. Kocsch, A Systematic Study of Fundamentals in α -Helical Coiled Coil Mimicry by Alternating Sequences of β - and γ -Amino Acids, *Amino Acids*, 2011, **41**, 733–742.
- 32 G. M. Whitesides, J. P. Mathias and C. T. Seto, Molecular Self-Assembly and Nanochemistry: a Chemical Strategy for



- the Synthesis of Nanostructures, *Science*, 1991, **254**, 1312–1319.
- 33 F. Thomas, W. M. Dawson, E. J. M. Lang, A. J. Burton, G. J. Bartlett, G. G. Rhys, A. J. Mulholland and D. N. Woolfson, De Novo-Designed α -Helical Barrels as Receptors for Small Molecules, *ACS Synth. Biol.*, 2018, **7**, 1808–1816.
- 34 F. Lapenta, J. Aupic, Z. Strmsek and R. Jerala, Coiled Coil Protein Origami: from Modular Design Principles towards Biotechnological Applications, *Chem. Soc. Rev.*, 2018, **47**, 3530–3542.
- 35 J. M. Fletcher, A. L. Boyle, M. Bruning, G. J. Bartlett, T. L. Vincent, N. R. Zaccai, C. T. Armstrong, E. H. Bromley, P. J. Booth, R. L. Brady, A. R. Thomson and D. N. Woolfson, A Basis Set of de Novo Coiled-Coil Peptide Oligomers for Rational Protein Design and Synthetic Biology, *ACS Synth. Biol.*, 2012, **1**, 240–250.
- 36 C. Toniolo, A. Polese, F. Formaggio, M. Crisma and J. Kamphuis, Circular Dichroism Spectrum of a Peptide 310-Helix, *J. Am. Chem. Soc.*, 1996, **118**, 2744–2745.
- 37 P. Schuck, On the Analysis of Protein Self-Association by Sedimentation Velocity Analytical Ultracentrifugation, *Anal. Biochem.*, 2003, **320**, 104–124.
- 38 K. Gade Malmos, L. Blancas-Mejia, B. Weber, J. Buchner, M. Ramirez-Alvarado, H. Naiki and D. Otzen, ThT 101: a Primer on the Use of Thioflavin T to Investigate Amyloid Formation, *Amyloid*, 2017, **24**, 1–16.
- 39 M. Biancalana and S. Koide, Molecular Mechanism of Thioflavin-T Binding to Amyloid Fibrils, *Biochim. Biophys. Acta, Proteins Proteomics*, 2010, **1804**, 1405–1412.
- 40 M. Zou, H. Yang, H. Wang, J. Zhang, B. Wei, H. Zhang and D. Xie, Detection of Type I Collagen Fibrils Formation and Dissociation by a Fluorescence Method Based on Thioflavin T, *Int. J. Biol. Macromol.*, 2016, **92**, 1175–1182.
- 41 E. K. Kumar, N. Hague and N. P. Prabhu, Kinetics of Protein Fibril Formation: Methods and Mechanisms, *Int. J. Biol. Macromol.*, 2017, **100**, 3–10.
- 42 R. Khurana, C. Coleman, V. Krishna, R. Roy and S. Singh, Mechanism of Thioflavin T Binding to Amyloid Fibrils, *J. Struct. Biol.*, 2005, **151**, 229–238.
- 43 K. Morimoto, K. Kawabata, S. Kunii, K. Hamano, T. Saito and B. Tonomura, Characterization of Type I Collagen Fibril Formation Using Thioflavin T Fluorescent Dye, *J. Biochem.*, 2009, **145**, 677–684.
- 44 M. Yan and X. Wang, Study on the Kinetic Self-Assembly of Type I Collagen from Tilapia (*Oreochromis Niloticus*) Skin Using the Fluorescence Probe Thioflavin T, *Spectrochim. Acta, Part A*, 2018, **203**, 342–347.
- 45 A. Adochitei and G. Drochioiu, Rapid Characterization of Peptide Secondary Structure by FT-IR Spectroscopy, *Rev. Roum. Chim.*, 2011, **56**, 783–791.
- 46 S. Spassov, M. Beekes and D. Naumann, Structural Differences between TSEs Strains Investigated by FT-IR Spectroscopy, *Biochim. Biophys. Acta*, 2006, **1760**, 1138–1149.
- 47 V. Serrano, W. Liu and S. Franzen, An Infrared Spectroscopic Study of the Conformational Transition of Elastin-Like Polypeptides, *Biophys. J.*, 2007, **93**, 2429–2435.
- 48 R. I. Litvinov, D. A. Faizullin, Y. F. Zuev and J. W. Weisel, The α -Helix to β -Sheet Transition in Stretched and Compressed Hydrated Fibrin Clots, *Biophys. J.*, 2012, **103**, 1020–1027.
- 49 V. A. Lorenz-Fonfria and E. Padros, Curve-Fitting of Fourier Manipulated Spectra Comprising Apodization, Smoothing, Derivation and Deconvolution, *Spectrochim. Acta, Part A*, 2004, **60**, 2703–2710.
- 50 R. Sarroukh, E. Goormaghtigh, J. M. Ruysschaert and V. Raussens, ATR-FTIR: A ‘Rejuvenated’ Tool to Investigate Amyloid Proteins, *Biochim. Biophys. Acta, Biomembr.*, 2013, **1828**, 2328–2338.
- 51 N. Kuhar, S. Sil, T. Verma and S. Umamathy, Challenges in Application of Raman Spectroscopy to Biology and Materials, *RSC Adv.*, 2018, **8**, 25888–25908.
- 52 A. Barth, Infrared Spectroscopy of Proteins, *Biochim. Biophys. Acta, Biomembr.*, 2007, **1767**, 1073–1101.
- 53 T. Miura, H. Takeuchi and I. Harada, Characterization of Individual Tryptophan Side Chains in Proteins Using Raman Spectroscopy and Hydrogen-Deuterium Exchange Kinetics, *Biochemistry*, 1988, **27**, 88–94.
- 54 R. W. Newberry, S. J. Orke and R. T. Raines, $n \rightarrow \pi^*$ Interactions Are Competitive with Hydrogen Bonds, *Org. Lett.*, 2016, **18**, 3614–3617.
- 55 C. Mensch, P. Bultinck and C. Johannessen, The Effect of Protein Backbone Hydration on the Amide Vibrations in Raman and Raman Optical Activity Spectra, *Phys. Chem. Chem. Phys.*, 2019, **21**, 1988–2005.
- 56 J. Seo, W. Hoffmann, S. Warnke, X. Huang, S. Gewinner, W. Schöllkopf, M. T. Bowers, G. von Helden and K. Pagel, An Infrared Spectroscopy Approach to Follow β -Sheet Formation in Peptide Amyloid Assemblies, *Nat. Chem.*, 2017, **9**, 39–44.
- 57 S. D. Moran and M. T. Zanni, How to Get Insight into Amyloid Structure and Formation from Infrared Spectroscopy, *J. Phys. Chem. Lett.*, 2014, **5**, 1984–1993.
- 58 E. Tayeb-Fligelman, O. Tabachnikov, A. Moshe, O. Goldshmidt-Tran, M. R. Sawaya, N. Coquelle, J. P. Colletier and M. Landau, The Cytotoxic Staphylococcus Aureus PSM α 3 Reveals a Cross- α Amyloid-Like Fibril, *Science*, 2017, **355**, 831–833.
- 59 S. Q. Zhang, H. Huang, J. Yang, H. T. Kratochvil, M. Lolicato, Y. Liu, X. Shu, L. Liu and W. F. DeGrado, Designed Peptides that Assemble into Cross- α Amyloid-Like Structures, *Nat. Chem. Biol.*, 2018, **14**, 870–875.
- 60 L. Lutter, C. J. Serpell, M. F. Tuite and W.-F. Xue, The Molecular Lifecycle of Amyloid – Mechanism of Assembly, Mesoscopic Organisation, Polymorphism, Suprastructures, and Biological Consequences, *Biochim. Biophys. Acta, Proteins Proteomics*, 2019, **1867**, 140257.
- 61 W. Lee, H. Lee, G. Lee and D. S. Yoon, Advances in AFM Imaging Applications for Characterizing the Biophysical



- Properties of Amyloid Fibrils, in *Exploring New Findings on Amyloidosis*, InTech, 2016.
- 62 D. Papapostolou, A. M. Smith, E. D. Atkins, S. J. Oliver, M. G. Ryadnov, L. C. Serpell and D. N. Woolfson, Engineering nanoscale order into a designed protein fiber, *Proc. Natl. Acad. Sci. U. S. A.*, 2007, **104**, 10853–10858.
- 63 T. H. Sharp, M. Bruning, J. Mantell, R. B. Sessions, A. R. Thomson, N. R. Zaccai, R. L. Brady, P. Verkade and D. N. Woolfson, (2012). Cryo-transmission electron microscopy structure of a gigadalton peptide fiber of *de novo* design, *Proc. Natl. Acad. Sci. U. S. A.*, 2012, **109**, 13266–13271.
- 64 O. S. Makin and L. C. Serpell, Structures for amyloid fibrils, *FEBS J.*, 2005, **272**, 5950–5961.
- 65 E. Tayeb-Fligelman, N. Salinas, O. Tabachnikov and M. Landau, *Staphylococcus aureus* PSM α 3 Cross- α Fibril Polymorphism and Determinants of Cytotoxicity, *Structure*, 2020, **28**, 301–313.
- 66 A. Lupas, M. Van Dyke and J. Stock, Predicting Coiled Coils from Protein Sequences, *Science*, 1991, **252**, 1162–1164.
- 67 E. Gasteiger, C. Hoogland, A. Gattiker, S. Duvaud, M. R. Wilkins, R. D. Appel and A. Bairoch, Protein Identification and Analysis Tools on the ExPASy Server, in *The Proteomics Protocols Handbook*, Human Press, 2005, pp. 571–607.
- 68 J. Dam and P. Schuck, Calculating Sedimentation Coefficient Distributions by Direct Modeling of Sedimentation Velocity Concentration Profiles, *Methods Enzymol.*, 2004, **384**, 185–212.
- 69 P. Schuck, Size-Distribution Analysis of Macromolecules by Sedimentation Velocity Ultracentrifugation and Lamm Equation Modeling, *Biophys. J.*, 2000, **78**, 1606–1619.
- 70 D. Hayes, T. Laue and J. Philo, *Program Sednterp: Sedimentation Interpretation Program*, Alliance Protein Laboratories, Thousand Oaks, CA, 1995.
- 71 H. Durchschlag and P. Zipper, Calculation of the Partial Volume of Organic Compounds and Polymers, *Prog. Colloid Polym. Sci.*, 1994, **94**, 20–39.
- 72 J. Vistica, J. Dam, A. Balbo, E. Yikilmaz, R. A. Mariuzza, T. A. Rouault and P. Schuck, Sedimentation Equilibrium Analysis of Protein Interactions with Global Implicit Mass Conservation Constraints and Systematic Noise Decomposition, *Anal. Biochem.*, 2004, **326**, 234–256.
- 73 P. Schuck, H. Zhao, C. A. Brautigam and R. Ghirlando, *Basic Principles of Analytical Ultracentrifugation*, CRC Press, 2016.
- 74 D. F. Kreitler, D. E. Mortenson, K. T. Forest and S. H. Gellman, Effects of Single α -to- β Residue Replacements on Structure and Stability in a Small Protein: Insights from Quasiracemic Crystallization, *J. Am. Chem. Soc.*, 2016, **138**, 6498–6505.
- 75 C. D. Schwieters, J. J. Kuszewski and G. M. Clore, *Prog. Nucl. Magn. Reson. Spectrosc.*, 2006, **48**, 47–62.
- 76 T. C. Huang, H. Toraya, T. N. Blanton and Y. Wu, X-ray powder diffraction analysis of silver behenate, a possible low-angle diffraction standard, *J. Appl. Crystallogr.*, 1993, **26**(2), 180–184.
- 77 R. Girardot, G. Viguier, M. Ounsy and J. Pérez, Foxtrot, 2010–2017, <https://www.synchrotron-soleil.fr/en/beamlines/swing>.

