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Stapling of leu-enkephalin analogs with bifunctional reagents for prolonged analgesic activity†

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The design and synthesis of leu-enkephalin analogs by replacing the glycine residues with *N*-(2-thioethyl)glycines and opening the cyclisation potential is presented. The cyclization (stapling) was achieved using bifunctional reagents (hexafluorobenzene and trithiocyanuric acid derivatives). The CD conformational studies of the stapled analogs suggest that the peptides adopt the type I β -turn conformation, which is in agreement with the theoretical analysis. The analog containing a trithiocyanuric acid derivative with a benzyl substituent shows potent analgesic activity.

In 1975, Hughes *et al.* discovered the endogenous opioid pentapeptides leu-enkephalin and met-enkephalin, which, in addition to their antinociceptive effects, have many other biological activities.¹ Structure–function studies have shown that two aromatic residues, Tyr and Phe, play a key role in the interaction with opioid receptors.^{2,3} This recognition is closely related to conformational preferences, which have been studied in detail, showing that the μ receptor requires the two rings to be on opposite sides of the peptide backbone, whereas δ receptors prefer aromatic residues on the same side.⁴ The structures of the enkephalins have been studied by X-ray crystallography, spectroscopic methods, and molecular modelling.^{5–9} These reports indicate that enkephalins are flexible and occur mainly in extended⁷ and folded conformations.^{6,8,9} A single turn stabilized by a β -turn located in the peptide region Gly²–Leu⁵ and a double turn in which the γ -turn is located on Gly² and the β -turn is centred on Gly³–Phe⁴, depending on the biomimetic environment used, are prominent folded conformations. Enkephalins' flexibility makes finding their active conformation tough. Studies suggest that in

opioid peptides, the residues in turn position not only restrict conformation but also directly interact with receptors.

The use of enkephalins in the treatment of pain is limited by their low metabolic stability and bioavailability, and their inability to cross the blood–brain barrier (BBB). Moreover, flexibility of enkephalins might cause side effects by binding to various receptors.¹⁰ The design of new analogs was aided by molecular modelling and receptor structure determination.^{11–13} Therefore, the synthesis of conformationally constrained leu-enkephalin analogs has been developed over many decades in the search for new peptidomimetics with improved stability and biological activity. Numerous analogs were obtained by cyclization,¹⁴ substitution of single amino acid residues with natural or unnatural amino acids,¹⁵ incorporation of cyclopropane-based scaffolds,¹⁶ introduction of linear and oligoheterocyclic motifs,¹⁷ introduction of amide bond isosters (ester, *N*-methylamide, triazole, alkene, trifluoroethylamine, azapeptide and fluoroalkene),¹⁸ introduction of sugar moieties,¹⁹ β -turn mimetic synthesis²⁰ and retropeptide synthesis.²¹

Stapling has been used to cross-link the peptide chains between *i*, *i* + 4 and *i*, *i* + 7 residues, stabilizing the secondary, mostly helical structure.²² In our approach, two glycine residues, placed in positions *i* and *i* + 1, were replaced by *N*-(2-thioethyl)glycine derivatives resulting in the leu-enkephalin peptoid (**1**) which was further cyclized through reactive –SH groups. The analog **1**, which was synthesized based on the modified protocol developed by our group,²³ was subjected to a cyclization reaction using two different reagents: hexafluorobenzene (Scheme 1a) and the trithiocyanuric acid derivative (Scheme 1b). As reported previously, cyclization through SH groups resulted in promising cyclic opioid peptides – DPDPE ((*D*-Pen2,*D*-Pen5)-enkephalin) in which the Gly² and Leu⁵ residues were replaced by the unnatural amino acid – Pen (penicillamine).^{14c} Details of our synthesis are provided in the ESI† (S2.1 and S2.3). The cyclization reagents were chosen to investigate different geometries of the target product, since hexafluorobenzene prefers 1,4-substitution,²⁴ while trithiocyanuric acid derivative prefers 1,3-substitution.²⁵ Hexafluorobenzene stapling

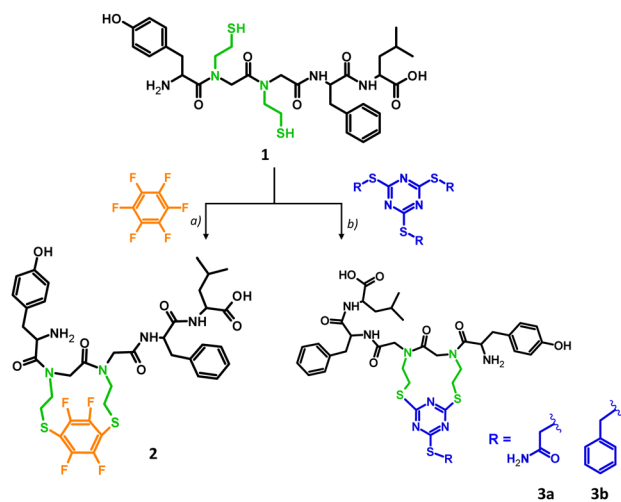
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Scheme 1 Synthesis scheme of the stapled analogs of leu-enkephalin (reaction conditions: (a) C_6F_6 (25 eq.), TCEP (5 eq.), 50 mM TRIS in DMF; (b) TMT(R)₃ (2 eq.), TCEP (5 eq.), 0.5 M TEAB (pH = 8.5) in DMF).

has so far been applied to -SH groups located in the side chain of amino acid residues at positions *i* and *i* + 4 of the peptide chain, resulting in a stable helical structure.²⁴ The stapling strategy based on sulfhydryl groups catching with 2-alkylsulfanyl-1,3,5-triazin-4,6-diyl linkers has been applied for the model peptide by the method inspired by recently published metathesis reactions of tris(alkyl) thiocyanurates.²⁵ These chemoselective reactions, occurring under mild basic conditions, are promising in designing agents for site-selective peptide or protein modification. Product **1** was characterized by LC-MS/MS and LC-UV. The LC-MS spectrum showed an expected signal at *m/z* 676.2865, corresponding to the molecular formula $C_{32}H_{45}N_5O_7S_2$ (Fig. S3A–C and S4, ESI[†]), which was confirmed by MS/MS *via* peptide sequence fragment ions (Fig. S3D, ESI[†]). The hexafluorobenzene reaction adapted from Spokoiny *et al.*²⁴ was modified by the introduction of a reducing reagent (TCEP) due to rapid intermolecular oxidation in the presence of TRIS, resulting in 65% yield of the product (Fig. S5, ESI[†]). Attempts at intramolecular disulphide bridge formation led to the formation of the dimeric product **1a** from the linear precursor with -SH groups. The detailed analysis of the dimer structure (parallel and antiparallel) was not continued. Stapled analogue **2** was purified by HPLC. The homogeneity and identity of the product were confirmed by LC-MS/MS (Fig. S6, ESI[†]), LC-UV (Fig. S15, ESI[†]) and NMR (Fig. S34–S39, ESI[†]) and the LC-MS spectrum showed a strong signal at *m/z* 822.2703 corresponding to the desired product with molecular formula $C_{38}H_{43}N_5O_7S_2F_4$ (Fig. S7, ESI[†]). Tandem mass spectrometry confirmed cyclic product formation, with identified ions derived from the linear portion of the peptide chain (Fig. S5E, ESI[†]). The expected 1,4 substitution of hexafluorobenzene²⁶ was confirmed by a single resonance registered at -73.3 ppm in the ¹⁹F NMR and consistent with a high symmetry (Fig. S39, ESI[†]). The cyclization towards **3a** was achieved with 2 eq. of TMT(Acm)₃ in 0.5 M TEAB H₂O buffer and a 1 : 1 mixture of DMF/20 mM TCEP (see the ESI[†], S2.1.5). The reaction is immediate and completes within 3 hours to give **3a** quantitatively as documented in the LC-MS analysis. The reference

3b derivative showed a limited solubility causing a need for extended reaction time and a full conversion after 24 hours of incubation (40 °C). The identities of both stapled analogues (**3a** and **3b**) were confirmed by ESI-MS and MS/MS analysis (Fig. S10–S13, ESI[†]). Products were purified chromatographically and the purity was confirmed by LC-UV analysis (Fig. S17 and S18, ESI[†]). CID fragmentation spectra of triazinyl analogs (**3a** and **3b**, Fig. S13, ESI[†]) are more complicated than those of the 4FB counterpart (**2**, Fig. S5E, ESI[†]). The most intense signals correspond to fragments derived by cleavage of bonds in the linear part. However, we also identified ions created by direct fragmentation of the linker or β elimination in one of stapled *N*-(2-SET)Gly residues followed by further fragmentation of the thiocyanurate moiety. The ¹H NMR spectrum of **2** and **3a** is complex due to presence of all four possible isomers (*cis-trans* isomerization of two tertiary amide bonds). In addition, DFT analysis of four possible isomers of analogs **2** and **3a** showed slight energy differences, confirming that all forms are equally probable. ¹H NMR comparative analysis of analogues **2** and **3a** (Fig. S42 and S43, ESI[†]) in most informative regions (amide, aromatic, aliphatic – Leu) shows clearly the existence of 2–4 forms. Well documented isomerization of peptidyl-proline²⁷ or peptoid²⁸ amide bonds allows conclusion of isomerization in the case of acyl-*N*-(2-mercaptoethyl)glycine, which fit to the observed spectra. We observed an increase in the number of forms in the triazine-stapled analogue. Analysis of the Tyr residue's -OH group (9.4–9.3 ppm) shows signal splitting into two for analogue **2** and into four for analogue **3a** (Fig. S43A, ESI[†]). The fluorobenzene-stapled peptide exhibits doubled para signals of tyrosine in the aromatic region (6.5–7 ppm), likely due to isomerization of the tyrosyl-*N*-(2-SET)Gly bond (Fig. S43C, ESI[†]). In the triazine-stapled analogue, a more conformationally labile molecule is indicated by a complex multiplet at 6.7 ppm and partially separated doublets from 6.95–7.15 ppm (two in each range: 6.9–7 ppm and 7.05–7.15 ppm). Similar complexity is observed for the -CH₃ groups of the leucine residue, identified as a doublet of doublets in Leu-enkephalin and a complex multiplet in stapled analogs. VT NMR analysis shows a temperature-dependent equilibrium in *cis-trans* isomerization of cyclic systems (Fig. S44–S46, ESI[†]).

The CD conformational studies were performed for leu-enkephalin, stapled analogues (**2**, **3a**, and **3b**), and the side product **1a** (dimer) in the far-UV region in three different solvents: MeCN, TFE, and H₂O (Fig. S19–S23, ESI[†]). The aromatic side chains of Tyr¹, Phe⁴ and bifunctional reagents used for stapling in the CD contributions obscure the peptide chromophore bands, making it hard to draw conformational conclusions. The CD spectra of **3a** and **3b** in TFE are quite similar to each other. They show negative bands at 250–260 nm, negative shoulders at 220–225 nm, and negative bands at 207 nm. The spectrum of **2** is somewhat similar to the two above ones below 220 nm, whereas at the longer wavelengths, there are only positive ellipticities, contrary to **3a** and **3b**. The CD spectra of all the analogues suggest that the peptides adopt the type I β -turn conformation in TFE. It was found that the CD spectrum typical of a type I β -turn is helical²⁹ and follows two CD patterns – one with a negative band at 224 nm, a negative



shoulder at 210 nm, and a positive band at 193 nm, and another with a negative shoulder at 224 nm, a negative band at 205 nm, and a positive band at 193 nm.³⁰ The conclusion concerning a type I β -turn in the analogues studied is consistent with the results of the theoretical calculations (Table S10, ESI†). The Φ and Ψ angle values found for Gly² and Gly³ residues of **2**, **3a**, and **3b** in the low-energy conformers correspond to the standard values of the type I β -turn.³¹ This specific type of β -turn has been found for leu-enkephalin by the NMR studies in DMSO-*d*₆.³² Moreover, also found in all the analogues were the intramolecular 4 \rightarrow 1 hydrogen bonds, between N-H of Phe⁴ and C=O of Tyr¹, stabilizing the β -turn conformation. The calculations showed also another conformational feature of **2**, **3a**, and **3b**, namely the presence of a γ -turn on the Phe⁴ residue, stabilized by the 3 \rightarrow 1 hydrogen bond, between Leu⁵ N-H and Gly³ C=O. A γ -turn gives a positive CD band at 230 nm.³³ Such a band can be seen in the spectrum of **2** but not in the spectra of **3a** and **3b**. In the latter two peptides, this band is probably overlapped by neighbouring negative ones. The CD spectra of **3b** in MeCN and water are similar to those in TFE, except for the longwave negative band. It suggests that in these two solvents, the conformation of **3b** is the same as in TFE (Fig. S23, ESI†).

In contrast, the spectra of **3a** are more difficult to interpret in MeCN and water, suggesting that the peptide experiences greater conformational flexibility in these solvents. Yet another situation is observed in the case of **2**. While one can suppose, on the basis of the CD spectra in the region of 200–220 nm, that the conformation of this analogue is somehow similar to that in TFE, in MeCN it seems to be distinctly different. The spectrum of **2** in MeCN in the far-UV region is more similar to the spectrum of leu-enkephalin in the same solvent rather than to spectra in TFE and water. As confirmed from the CD spectra, the presented solvent dependence of the tested analogues shows that the introduced bridge on the adjacent substituted glycine residues still leaves the conformational freedom of the entire system, but clearly affects the secondary structure. Computational methods have been used as a useful tool to predict the structure of the molecules, ligands and complexes.³⁴ Molecular orbital studies on **2**, **3a**, **3b** and **Leu-Enk** have been done at the DFT level of theory with the IEFPCM (integral equation formalism for polarizable continuum model) solvent (water) approach. The starting structure of the peptides for DFT calculations was generated on the basis of the amino acid sequence after 45 ps simulation at 300 K, without cutoffs using BIO+ implementation of the CHARMM force field. The results of the conformational studies correlate very well with the DFT analysis of the stapled analogues. At the DFT level of theory, we have found four thermodynamically stable conformations **2**, **3a**, **3b** and **Leu-Enk** displaying terminus-terminus interaction with PA \cdots PD distances 2.649 Å, 2.532 Å, 2.534 Å and 2.668 Å respectively (Fig. S47–S50, ESI†). Despite their limited size, these molecules form consistent intramolecular hydrogen bonds, as shown in Table S11 (ESI†). The structures of stapled **2**, **3a** and **3b** are stabilized by hydrogen bonds between Tyr¹ and Phe⁴ residues resulting in a 10-member ring with values 2.879 Å,

2.974 Å and 2.794 Å, respectively. An additional gamma-bend stabilized by hydrogen bonds (2.885 Å and 2.888 Å) between *N*-(2-Set)Gly³ and Leu⁵ is observed in the triazine analogues (**3a** and **3b**). The reference compound (**Leu-Enk**) displays a 7-member HB bonded ring formed between Tyr¹ and Gly² with 2.878 Å PA \cdots PD distance as presented in Fig. S49 (ESI†). The distance between the most remote carbon atoms in the aromatic rings of **3b** is \sim 20 Å, while that in **3a** is only \sim 10 Å. The cartesian coordinates of presented molecules can be found in the ESI† (Table S12).

Biological tests involved two stapled analogues (**2** and **3a**) and unaltered leu-enkephalin as the reference compound. An analogue featuring a triethiocyanuric acid derivative was tested alongside another with an added benzyl system. Leu-enkephalin was synthesized following the standard Fmoc protocol; analytical data are provided in the ESI.† Antinociceptive activity of analogues **2** and **3b** was assessed in Wistar rats *via* intrathecal administration (i.t.) using a tail flick test, and the results are shown in Fig. 1. All analogues displayed time-dependent analgesic effects at 16 nmol kg^{−1} i.t. dose, with distinct differences between **2** and **3b**. Analog **2** peaked at 30 min, exhibiting a comparable analgesic effect to **3b** (\sim 50% MPE) and leu-enkephalin. Its lowest efficacy (\sim 20% MPE) was observed at 120 min.

A different profile of analgesic activity was shown by **3b**. The lowest activity was at 5 min (20.72% MPE); at 30 min the effect was similar to that of **2**, and the highest effect was at 120 min, which was higher than that of Leu-Enkephalin (20.72% MPE to 65.2% MPE). At 120 min post-administration, **2** exhibited weak activity where **3b** reached the highest peak suggesting this compound may have longer duration of action than 120 min. Overall, the antinociceptive action of the tested analogues was relatively high, comparable to the standard leu-enkephalin. It should be noted that **3b** has other analgesic profile response than **2** and leu-enkephalin. Its activity increased in the following minutes to reach its maximum at 120 min. The effect of the obtained derivatives (**2** and **3b**) is analogous to that of the action of leu-enkephalin, as demonstrated by analgesic tests. The prolonged action of the stapled analog can be explained by higher enzymatic stability for cyclic peptides. This observation requires further investigation because it may be important for

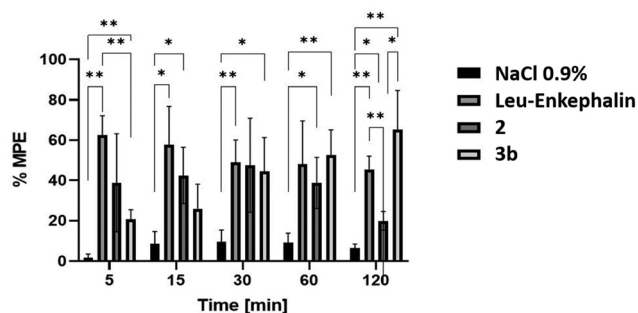


Fig. 1 Analgesic responses of analogs **2** and **3b** as well as **Leu-Enk** (standard). Results were analyzed using two-way repeated measures of ANOVA followed by Tukey's multiple comparison test and plotted as the mean \pm SEM ($n = 7-8$). Asterisks (*) indicate a significant difference $*p < 0.03$, $**p < 0.001$.



the development of tolerance with chronic administration for this compound.

Biological studies confirmed the activity of the stapled analogs, suggesting that their rigid structure interacts effectively with the receptor. Identification of the bioactive conformer remains challenging due to isomerization induced by tertiary amide bonds observed by NMR analysis. Theoretical and CD analyses show structural similarities among the analogs, suggesting the importance of aromatic ring orientation. However, the determination of the privileged structural motif for effective receptor interaction remains elusive. The presence and orientation of the additional benzyl substituent likely enhance the prolonged analgesic effect of analog **3b**.

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Conflicts of interest

There are no conflicts to declare.

Notes and references

- (a) P. W. Schiller, *Prog. Med. Chem.*, 1991, **28**, 301–340; (b) J. E. Holden, Y. Jeong and J. M. Forrest, *AACN Adv. Crit. Care*, 2005, **16**, 291–301; (c) M. S. Henry, L. Gendron, M.-E. Tremblay and G. Drolet, *Neural Plast.*, 2017, 1546125; (d) I. I. Shcheniavsky, *Probl. Cryobiol. Cryomed.*, 2021, **31**, 003–013; (e) C. V. Borlongan, Y. Wang and T. P. Su, *Front Biosci.*, 2004, **9**, 3392–3398.
- V. J. Hruby and C. A. Gerhring, *Med. Res. Rev.*, 1989, **9**, 343–401.
- (a) H. Kimura, C. H. Stammer, Y. Shimohigashi, C. Ren-Lin and J. Stewart, *Biochem. Biophys. Res. Commun.*, 1983, **115**, 112–115; (b) Y. Shimohigashi, T. Costa, A. Pfeiffer, A. Herz, H. Kimura and C. H. Stammer, *FEBS Lett.*, 1987, **222**, 71–74.
- D. Tourwé, K. Verschuere, A. Frycia, P. Davis, F. Porreca, V. J. Hruby, G. Toth, H. Jaspers, P. Verheyden and G. Van Binst, *Biopolymers*, 1996, **38**, 1–12.
- D. Mastropalo, A. Camerman, L. Y. Y. Ma and N. Camerman, *Life Sci.*, 1987, **40**, 1995–1999.
- C. Garbay-Jauregui, B. P. Roques, R. Oberlin, M. Anteunis, S. Combrisson and J. Y. Lallemand, *FEBS Lett.*, 1977, **76**, 93–98.
- L. Rudolph-Bohner, D. Quarzago, M. Czisch, U. Ragnarsson and S. Moroder, *Biopolymers*, 1997, **41**, 591–606.
- D. Picone, A. D'Ursi, A. Motta, T. Tancredi and P. A. Temussi, *Eur. J. Biochem.*, 1990, **192**, 433–439.
- A. Milon, T. Miyazawa and T. Higashijima, *Biochemistry*, 1990, **29**, 65–75.
- A. Janecka, J. Fichna and T. Janicki, *Curr. Top. Med. Chem.*, 2004, **4**, 1–17.
- I. Efremenko and R. H. Fish, *Organometallics*, 2015, **34**, 4117–4126.
- H. S. Park, B. J. Byun and Y. K. Kang, *ACS Omega*, 2022, **7**(31), 27755–27768.
- F. Wieberneit, A. Korste, H. B. Albada, N. Metzler-Nolte and R. Stoll, *Dalton Trans.*, 2013, **42**, 9799.
- (a) G. Toth, T. H. Kramer, R. Knapp, G. Lui, P. Davis, T. F. Burks, H. I. Yamamura and V. J. Hruby, *J. Med. Chem.*, 1990, **33**, 249–253; (b) M. H. Hannah, O. E. Shainnel, M. L. Ganno, J. C. Davis, C. T. Dooley, J. P. McLaughlin and A. Nefzi, *Org. Biomol. Chem.*, 2019, **17**, 5305–5315; (c) G. Toth, T. H. Kramer, R. Knapp, G. Lui, P. Davis, T. F. Burks, H. I. Yamamura and V. J. Hruby, *J. Med. Chem.*, 1990, **33**, 249–253.
- (a) H. Xue, M. Guo, Ch Wang, Y. Shen, R. Qi, Y. Wu, Z. Xu and M. Chang, *Org. Chem. Front.*, 2020, **7**, 2426–2431; (b) S. Maricic, U. Berg and T. Frejd, *Tetrahedron*, 2002, **58**, 3085–3093; (c) Y. Sasaki, M. Hirabuki, A. Ambo, H. Ouchi and Y. Yamamoto, *Bioorg. Med. Chem. Lett.*, 2001, **11**, 327–329; (d) H. Xue, M. Guo, Ch Wang, Y. Shen, R. Qi, Y. Wu, Z. Xu and M. Chang, *Org. Chem. Front.*, 2020, **7**, 2426–2431.
- A. Mizuno, K. Matsui and S. Shuto, *Chem. – Eur. J.*, 2017, **23**, 14394–14409.
- S. Hammami, Z. Mighri, C. T. Dooley and A. Nefzi, *Bioorg. Med. Chem. Lett.*, 2014, **24**, 4482–4485.
- (a) K. Rochon, A. Proteau-Gagne, P. Bourassa, J.-F. Nadon, J. Cote, V. Bournival, F. Gobeil, Jr., B. Guerin, Y. L. Dory and L. Gendron, *ACS Chem. Neurosci.*, 2013, **4**, 1204–1216; (b) S. Sun, Q. Jia and Z. Zhang, *Bioorg. Med. Chem. Lett.*, 2019, **29**(18), 2535–2550; (c) J.-L. Beaudreau, V. Blais, B. J. Holleran, A. Bergeron, G. Piñeyro, B. Guerin, L. Gendron and Y. L. Dory, *ACS Chem. Neurosci.*, 2019, **10**, 1615–1626; (d) S. N. Karad, M. Pal, R. S. Crowley, T. E. Prisinzano and R. A. Altman, *Chem. Med. Chem.*, 2017, **12**, 571–576; (e) R. Sinisi, A. Ghilardi, S. Rui, P. Lazzari, L. Malpezzi, M. Sani, L. Pani and M. Zanda, *Chem. Med. Chem.*, 2009, **4**, 1416–1420; (f) V. Eeda, M. Selvaraju and R. A. Altman, *J. Fluorine Chem.*, 2019, **218**, 90–98; (g) M. O. Bowles and C. Proulx, *Org. Lett.*, 2022, **24**, 1768–1773.
- M. J. Slusarz, *Comput. Biol. Chem.*, 2022, **101**, 107783.
- C. Xiong, J. Zhang, P. Davis, W. Wang, J. Ying, F. Porreca and V. J. Hruby, *Chem. Commun.*, 2003, 1598–1599.
- N. Soumanou, D. Lybye, T. Hjelmgaard and S. Faure, *Green Chem.*, 2023, **25**, 3615–3623.
- (a) X. Li, S. Chen, W.-D. Zhang and H.-G. Hu, *Chem. Rev.*, 2020, **120**, 10079–10144; (b) M. Kijewska, G. Wolczański, M. Światowska, K. Kędziora, M. Pawlicki and P. Stefanowicz, *Chem. – Eur. J.*, 2023, **29**, e202301370; (c) M. Kijewska, A. Czerwińska, S. Al-Harthi, G. Wolczański, M. Waliczek, A.-H. Emwas, M. Jaremko, E. Jaremko, P. Stefanowicz and Z. Szewczuk, *Chem. Commun.*, 2020, **56**, 8814–8817.
- M. Wierzbicka, M. Waliczek, A. Dziadecka and P. Stefanowicz, *J. Org. Chem.*, 2021, **86**, 12292–12299.
- A. M. Spokoiny, Y. Zou, J. J. Ling, H. Yu, Y.-S. Lin and B. L. Pentelute, *J. Am. Chem. Soc.*, 2013, **135**(16), 5946–5949.
- G. Wolczański, W. Gil, J. Cichos, M. Lisowski and P. Stefanowicz, *J. Org. Chem.*, 2023, **88**, 8192–8202.
- P. Dognin, P. M. Killoran, G. S. Hanson, L. Halsall, T. Chaudhry, Z. Islam, F. Giuntini and Ch. R. Coxon, *Peptide Sci.*, 2021, **113**, e24182.
- U. Reimer, G. Scherer, M. Drewello, S. Kruber, M. Schutkowski and G. Fischer, *J. Mol. Biol.*, 1998, **279**, 449–460.
- D. Kalita, B. Sahariah, S. P. Mookerjee and B. K. Sarma, *Chem. – Asian J.*, 2022, **17**, e2022001.
- J. Bandekar, D. J. Evans, S. Krimm, S. J. Leach, S. Lee, J. R. McQuie, E. Minasian, G. Nemethy, M. S. Pottle, H. A. Scheraga, E. R. Stimson and R. W. Woody, *Int. J. Peptide Protein Res.*, 1982, **19**, 187–205.
- A. Perczel and G. D. Fasman, *Protein Sci.*, 1992, **1**, 378–395.
- M. Migliore, A. Bonvicini, V. Tognetti, L. Guilhaudis, M. Baaden, H. Oulyadi, L. Jouberta and I. Segalas-Milazzo, *Phys. Chem. Chem. Phys.*, 2020, **22**, 1611–1623.
- (a) C. Garbay-Jauregui, B. P. Roques, R. Oberlin, M. Anteunis, S. Combrisson and J. Y. Lallemand, *FEBS Lett.*, 1977, **76**, 93–98; (b) E. R. Stimson, Y. C. Meinwald and H. A. Scheraga, *Biochemistry*, 1979, **18**, 1661–1671.
- E. Vass, Z. Majer, K. Köhalmi and M. Hollósi, *Chirality*, 2010, **22**, 762–771.
- (a) Z. Mielke, Z. Latajka, A. Olbert-Majkut and R. Wiczorek, *J. Phys. Chem. A*, 2000, **104**, 3764–3769; (b) R. Wiczorek, Z. Latajka and L. Lundell, *J. Phys. Chem. A*, 1999, **103**, 6234–6239; (c) T. K. Olszewski, E. Wojaczyńska, R. Wiczorek and J. Bąkiewicz, *Tetrahedron: Asymmetry*, 2015, **26**, 601–607; (d) P. Salvador, R. Wiczorek and J. J. Dannenberg, *J. Phys. Chem. B*, 2007, **111**, 2398–2403; (e) M. Rudowska, R. Wiczorek, A. Kluczyk and P. Stefanowicz, *JASMS*, 2013, **24**, 846–856; (f) E. Gumienna-Kontecka, G. Berthon, I. O. Fritsky, R. Wiczorek, Z. Latajka and H. Kozłowski, *J. Chem. Soc., Dalton Trans.*, 2000, 4201–4208.

