Syntheses and Self-assembling Behaviors of Pentagonal Conjugates of Tryptophane Zipper-Forming Peptide

Kazunori Matsuura 1,2,* , Kazuya Murasato 1 and Nobuo Kimizuka 1,2

1 Department of Chemistry and Biochemistry, Graduate School of Engineering, Kyushu University, Moto-oka 744, Nishi-ku, Fukuoka 819-0395, Japan;
E-Mails: ma2ra-k@mail.cstm.kyushu-u.ac.jp (K.M.); n-kimi@mail.cstm.kyushu-u.ac.jp (N.K.)
2 International Research Center for Molecular Systems, Kyushu University, Moto-oka 744, Nishi-ku, Fukuoka 819-0395, Japan
* Author to whom correspondence should be addressed; E-Mail: ma2ra-k@mail.cstm.kyushu-u.ac.jp;
Tel.: +81-92-802-2833; Fax: +81-92-802-2838.

Abstract: Pentagonal conjugates of tryptophane zipper-forming peptide (CKTWTLWTE) with a pentaazacyclopentadecane core (Pentagonal-Gly-Trpzip and Pentagonal-Ala-Trpzip) were synthesized and their self-assembling behaviors were investigated in water. Pentagonal-Gly-Trpzip self-assembled into nanofibers with the width of about 5 nm in neutral water (pH 7) via formation of tryptophane zipper, which irreversibly converted to nanoribbons by heating. In contrast, Pentagonal-Ala-Trpzip formed irregular aggregates in water.

1. Introduction

Multivalent ligand–receptor interactions play pivotal roles in biological systems [1–4]. To date, many artificial multivalent bioconjugates have been developed as inhibitors, receptors, artificial enzymes, signaling molecules, and drug delivery materials. For example, inhibition of Shiga-like or cholera toxins by pentavalent conjugates of oligosaccharides has been reported [5,6].

The concept of “template-assembled synthetic proteins (TASP)” provided a chemical approach to design artificial proteins [7]. Peptide dendrimers have also been developed as multifunctional biomaterials [8,9].

Many artificial peptide nano-assemblies have been designed by self-assembly of coiled-coil α-helix peptides and β-sheet-forming peptides [10–21]. Application of multivalent molecular design to peptide assembly can promote self-assembly and lead to unique morphologies. For example, Ghosh et al. reported that PAMAM dendrimer containing four leucine zipper-forming peptides self-assembled into nanofibers [22]. We have designed trigonal peptide conjugates containing β-sheet forming peptides [23–25] and glutathione [26–28], and demonstrated that they are useful components for the spontaneous construction of peptide nanoarchitectures in water. Tryptophane zipper has been attracting much attention as a secondary structure motif, which forms stable twisted β-hairpin structure due to the interaction between tryptophane residues [29–34]. Recently, we have reported that a novel trigonal peptide conjugate bearing tryptophane zipper-forming peptides showed pH-responding self-assembly into nanospheres and nanofibers (Figure 1) [35].

Most spherical virus capsids are self-assembled from some multiple of 60 chemically identical protein subunits and have an icosahedral symmetry which possesses three- and five-fold rotation axes [36,37]. Molecular design of C5-symmetric self-assembling molecules would provide chemical strategy for artificial virus capsid, since three-dimensional tiling of pentagon affords dodecahedron. Olson et al. demonstrated by molecular dynamics that C5-symmetric corannulene-based molecules have the potential to self-assemble into dodecahedral nanocapsule [38]. In this paper, we designed a pentagonal peptide conjugate bearing tryptophane zipper-forming peptides (Figure 1), and envisioned that the pentagonal peptide conjugate self-assembles into nanocapsules by formation of intermolecular tryptophane zipper structures. 1,4,7,10,13-Pentaazacyclopentadecane scaffold [6,39] was adopted as a pentagonal core, since the syntheses of pentagonal derivatives are easy to design.
2. Results and Discussion

2.1. Synthesis of Pentagonal Conjugates of Tryptophane Zipper-Forming Peptide

1,4,7,10,13-Pentaazacyclopentadecane 1 was synthesized from triethylenetetramine according to the reported procedure [39]. 1 was amidated with Boc-Ala or Boc-Gly, followed by deprotection and iodoacetylation to provide pentagonal core 4 (Scheme 1). \(^1\)H NMR spectrum of Ala-containing pentagonal core 4a in mixture of CD\(_3\)CN/D\(_2\)O = 15/1 showed notable broad peaks at the range of \(\delta 4.2–3.0\) ppm assigned to pentaazacyclopentadecane (\(H_a\)) and multiple peaks at \(\delta 4.7\) (\(H_b\)), 3.7 (\(H_c\)) and 1.2 ppm (\(H_d\)) (Figure 2). The multiple peaks of \(H_c\) were unified by elevating temperature, whereas the peaks assigned to \(H_a\) were broad even at 60 °C. This indicates that Ala-containing pentagonal core 4a possesses various irregular conformations of which interconversion are very slow even at 60 °C. In contrast, \(^1\)H NMR spectrum of Gly-containing pentagonal core 4b showed relatively sharp peaks (Figure 2), suggesting that 4b possesses flexible conformation. It is probable that steric hindrance of methyl group of Ala to pentaazacyclopentadecane ring of 4a prevent flexible interconversion of conformers, whereas steric hindrance in 4b is smaller than that of 4a.

The 8-mer peptide H-CKTWTWTE-OH, which was designed based on the intermolecular tryptophane zipper (β-hairpin) forming peptide reported by Cohhan et al. [29], was synthesized by a standard Fmoc-protected solid-phase method. Pentagonal-Ala-Trpzip and Pentagonal-Gly-Trpzip were prepared by coupling the 8-mer peptides with pentagonal core 4 in the presence of diisopropylethylamine (Scheme 1). These pentagonal conjugates were purified by reverse-phase HPLC and confirmed by MALDI-TOF-MS (\(m/z\) of Pentagonal-Ala-Trpzip = 6038.96 [M + H]+, \(m/z\) of Pentagonal-Gly-Trpzip = 5970.58 [M + H]+). They were soluble in water at a whole range of pH.

2.2. Secondary Structure of Pentagonal Peptide Conjugates

After the solutions of pentagonal peptide conjugates were incubated for 48 h at 25 °C in 20 mM citrate buffer (pH 3), in 20 mM phosphate buffer (pH 7), and in aqueous NaOH solution (pH 11), circular dichroism (CD) spectra were measured [40]. It has been reported that tryptophane zipper-forming β-hairpin peptide showed exciton-coupling type CD spectrum (positive peak at 230 nm and negative peak at 215 nm) along with small peaks at 285–295 nm based on the interaction between Trp residues [33]. The CD spectra of aqueous solution of Pentagonal-Gly-Trpzip showed negative peak at 216 nm and positive peak at 233 nm together with weak peaks at 280–300 nm in phosphate buffer (pH 7) and aqueous NaOH solution (pH 11) respectively. They indicate the formation of tryptophane zipper and normal β-sheet structure, whereas the CD intensity at 216 and 233 nm considerably decreased in citrate buffer (pH 3) (Figure 3(a)). In contrast, the CD spectra of Pentagonal-Ala-Trpzip showed a weak CD pattern at pH 7 and pH 3 which can be ascribed to tryptophane zipper structure, and the content of random coil increased at pH 11 (Figure 3(b)). The difference in secondary structure between these pentagonal Trpzip conjugates might arise from different flexibility of the pentagonal core, as shown by the \(^1\)H NMR spectra (Figure 2).

The CD spectrum of the precursor peptide CKTWTWTE revealed that the peptide adopted random-coil structure at the pH range of 3–11 (Figure 3(c)). We have previously reported that Trigonal-Trpzip also adopts mixed secondary structures of tryptophane zipper and normal β-sheet at pH 7, but the molar ellipticity at 216 and 233 nm ([\(\theta\])\(_{216}\) = –55,000 and [\(\theta\])\(_{233}\) = 21,000 deg cm\(^2\) dmol\(^–1\)) was lower than that of Pentagonal-Gly-Trpzip under the same conditions [35]. These results indicate that the formation of tryptophane zipper structure from the peptide CKTWTWTE is promoted by the pentagonal preorganization, which is more effective than the trigonal preorganization.

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**Scheme 1. Synthesis of pentagonal conjugates of tryptophane zipper peptide.**
2.3. Self-Assembly of Pentagonal Peptide Conjugates in Water

A transmission electron microscopy (TEM) image revealed that Pentagonal-Gly-Trpzip formed only nanofibers of several micrometers length with uniform width of 4–5 nm in phosphate buffer (pH 7, Figure 4(a)), although we envisioned that the conjugate self-assembles into nanocapsules [40]. It is probable that the nanofibers formed by face-to-face assembly of Pentagonal-Gly-Trpzip via parallel tryptophane zipper and β-sheet structure. On the other hand, in the TEM image of Pentagonal-Ala-Trpzip, irregular aggregates were observed together with nanofibers (Figure 4(b)). This might reflect less formation of tryptophane zipper structure based on rigid irregular core conformations.

We have previously reported that Trigonal-Trpzip selectively self-assembled into nanospheres with the size of 20–30 nm at pH 7 [35]. Since the peptide CKTWTWTE might possess zwitterionic structure at pH 7, it is reasonable that Trigonal-Trpzip formed nanospheres by forming intermolecular antiparallel β-sheet-like structures. In contrast, Pentagonal-Gly-Trpzip selectively formed nanofibers probably due to the formation of parallel β-sheet-like structures despite ionic repulsion between peptides (Figure 5). To form nanospheres, it is desirable that the assembly units take concave conformations. The difference in morphology between trigonal- and pentagonal-tryptophanezipper conjugates might arise from difference in the peripheral density of peptide chains and the conformation of scaffold. It seems that steric hindrance among peptide chains in Pentagonal-Gly-Trpzip prevents the formation of antiparallel β-sheet-like structures and concave conformations, in contrast to the case of Trigonal-Trpzip.

Figure 6(a) shows temperature dependence of CD spectrum for aqueous solution of Pentagonal-Gly-Trpzip (10 µM) at pH 7. The positive peaks at 233 and 285–295 nm were gradually decreased by heating process, but the CD intensity at 233 nm hardly recovered by cooling process (Figure 6(b)). CD spectrum of Pentagonal-Gly-Trpzip at 25 °C after the heating process showed formation of normal β-sheet-rich structure (Figure 6(c)). These results indicate the irreversible structural change of tryptophane zipper to normal β-sheet-rich structure.

A TEM image revealed that Pentagonal-Gly-Trpzip formed left-handed helical ribbon structures with the width of 20–30 nm and pitch of 50–100 nm after the heating process (Figure 6(d)). It is probable that kinetically self-assembled nanofibers consisting of tryptophane zipper transformed to thermodynamically stable helical ribbons consisting of normal β-sheet by the heating process.

3. Experimental Section

3.1. General

Reagents were obtained from commercial source and used without further purification. Deionized water of high resistivity (>18 MΩ cm) purified with a Millipore Purification System (Milli-Q water) was used as a solvent of peptide conjugates. 1H-NMR spectra were recorded on Bruker AV300M spectrometer. Reversed-phase HPLC was performed at ambient temperature with a Simadzu LC-6AD liquid chromatograph equipped with a UV/Vis detector (220 nm, Shimadzu SPD-10AVvp) using Inertsil ODS-3 (GL Science) or COSMOSIL Protein-R (Nakarai Tesque) columns (250 × 4.6 mm or 250 × 20 mm). MALDI-TOF mass spectra were obtained on Autoflex III (Bruker Daltonics) under the linear/positive mode with α-cyano-4-hydroxy cinnamic acid (α-CHCA) as matrix.

Figure 2. 1H-NMR spectra of pentagonal iodoacetomidated core molecules (4a and 4b) at 2.8 mM in mixture of CD$_3$CN/D$_2$O = 15/1.

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3.2. Synthesis of Peptide Conjugates

1,4,7,10,13-Pentaazacyclopentadecane (1): 1 was synthesized from triethylenetetramine according to the reported procedure [39].

Tryptophene zipper-forming peptide (H-CKTWTWTE-OH): Peptide H-Cys(Trt)-Lys(Boc)-Thr(tBu)Trp(Boc)-Thr(tBu)-Trp(Boc)-Thr(tBu)-Glu(OtBu)-Alko resin was synthesized on α-p-alkoxybenzyl alcohol resin (Alko resin, Watanabe Chemical Industries, Ltd., Hiroshima, Japan, 0.69 mmol/g) using standard Fmoc-based FastMoc coupling chemistry (3 eq. Fmoc-amino acids) with an ABI 433A synthesizer (Applied Biosystems, Carlsbad, CA, USA). DMF solution of 2-([1H-benzotriazole-1-yl]1,1,3,3-tetramethyluronium hexafluorophosphate (HBTU, 0.5 M) and 1-hydroxybenzotriazole hydrate (HOBt·H2O, 0.5 M) was used as a coupling reagent. 2.0 M diisopropylethylamine (DIPEA) in NMP and 20% piperidine in NMP were used for neutralization and for Fmoc deprotection, respectively. The peptidyl-resin was washed with NMP, dichloromethane and methanol then dried under vacuum. The peptide was deprotected and cleaved from the resin by treatment with a cocktail of TFA/1,2-ethanediol/water/trisopropylsilane = 94/2.5/2.5/1 in volume at room temperature for 2 h. The reaction mixture was filtered to remove the resin and the filtrate was concentrated under vacuum. The peptide was precipitated by adding ice-cooled methyl-tert-butyl ether to the residue and the supernatant was decanted. After repeating the methyl-tert-butyl ether washing 6 times, the precipitated peptide was dried under vacuum. The crude product was purified by reversed-phase HPLC (column: Inertsil ODS-3) eluting with a linear gradient of CH3CN/water (18/82 to 30/70 over 60 min) containing 0.1% TFA. The elution fraction containing the desired peptide was lyophilized to give a flocculent solid. The isolated yield was 23%. MALDI-TOF-MS (matrix: α-CHCA): m/z = 1054.47 [M + H]+. 1H-NMR (D2O, δ/ppm): 7.37–7.45 (2 H, m), 7.23–7.28 (2H, m), 6.90–7.08 (6H, m), 4.51 (1H, t, J = 7.2 Hz), 4.45 (1H, t, J = 7.2 Hz), 4.20 (1H, t, J = 7.2 Hz), 4.15 (1H, d, J = 5.1 Hz), 4.04–4.10 (3H, m), 3.85–4.01 (4H, m), 3.10 (2H, t, J = 6.3 Hz), 2.89–2.97 (4H, m), 2.67 (2H, t, J = 7.8 Hz), 2.22 (2H, t, J = 7.8 Hz), 1.84–1.99 (1H, m), 1.68–1.84 (1H, m), 1.48–1.58 (2H, m), 1.37–1.48 (2H, m), 1.07–1.20 (2H, m), 0.98 (3H, d, J = 6.3 Hz), 0.91 (3H, d, J = 6.3 Hz).

**Figure 3.** Circular dichroism (CD) spectra of aqueous solution of (a) Pentagonal-Gly-Trpzip (10 μM); (b) Pentagonal-Ala-Trpzip (10 μM); and (c) tryptophane zipper peptide CKTWWTETE (50 μM) at 25 ºC in 20 mM citrate buffer (pH 3, red line), in 20 mM phosphate buffer (pH 7, green line), and in aqueous NaOH solution (pH 11, blue line).

**Figure 4.** Transmission electron microscopy (TEM) images of pentagonal conjugates of tryptophane zipper peptide (10 μM) in 20 mM phosphate buffer (pH 7) at 25 ºC: (a) Pentagonal-Gly-Trpzip, and (b) Pentagonal-Ala-Trpzip. TEM samples were stained with phosphotungstic acid.
N,N”,N’,N’’,N’’’,N’’’’-Pentakis(N-tert-butoxycarbonyl-L-alanyl)-1,4,7,10,13-pentaazacyclododecane (2a): Boc-Ala-0H (265 mg, 1.40 mmol) and O-(7-azabenzotriazolyl)-tetramethyluronium hexafluorophosphate (HATU, 558 mg, 1.43 mmol) were dissolved in dry DMF (2.0 mL). DIPEA (244 µL, 1.40 mmol) was added to the mixture by microsyringe, and then the mixture was stirred for 45 min at room temperature. Then a solution of 1,4,7,10,13-pentaazacyclododecane 1 (30 mg, 0.14 mmol) in dry DMF (1.0 mL) was added to the mixture. After the mixture was stirred for 43 h at room temperature, the solvent was evaporated under reduced pressure. The residue was dissolved in chloroform and washed with 5% aqueous NaHCO₃ and deionized water. The organic layer was dried over anhydrous Na₂SO₄ and evaporated to a dry residue. The residue was purified by reversed-phase HPLC (column: Inertsil ODS-3) eluting with a linear gradient of CH₃CN/water (30/70 to 50/50 over 20 min). The isolated yield was 12%. MALDI-TOF-MS (matrix: α-CHCA): m/z = 1409.88 [M + H]+. ¹H-NMR (CD₃CN + D₂O, δ/ppm): 2.6–2.7 (10H, br), 1.8–1.9 (10H, br), 0.7–1.1 (20H, br).

N,N”,N’,N’’,N’’’,N’’’’-Pentakis(N-iodoacetylglglycyl)-1,4,7,10,13-pentaazacyclododecane (2b): Compound 2b was prepared by the almost same procedure described above. The crude yield was 96% (135 mg). The product was not purified further and was used in the next reaction. MALDI-TOF-MS (matrix: α-CHCA): m/z = 1092.64 [M + Na]+. ¹H-NMR (CD₃OD, δ/ppm): 3.4–4.9 (25H, br), 1.48 (45H, br), 1.37 (15H, br).

N,N”,N’,N’’,N’’’,N’’’’-Pentakis(N-iodoacetyl-L-alanyl)-1,4,7,10,13-pentaazacyclododecane (2c): A solution of 2b (47 mg, 0.044 mmol) and NaHCO₃ (28 mg, 0.33 mmol) in deionized water (1.0 mL) was added to a solution of iodoacetic acid (96 mg, 0.34 mmol) in acetone (1.0 mL). The mixture was stirred for 1 h in the dark at room temperature. After the solvent was evaporated under reduced pressure, the residue was purified by reversed-phase HPLC (column: Inertsil ODS-3) eluting with a linear gradient of CH₃CN/water (25/75 to 50/50 over 25 min). The isolated yield was 13.4 mg (23%). MALDI-TOF-MS (matrix: α-CHCA): m/z = 1340.79 [M + H]+. ¹H-NMR (CD₃CN + D₂O, δ/ppm): 7.5–7.6 (2H, br), 4.10 (10H, s), 3.76 (10H, s), 3.4–3.6 (20H, br).

Pentagonal-Ala-Trpzip: Peptide H-CKTWTWTE-OH (13.8 mg, 10.8 µmol) was dissolved in degassed DMF (4.0 mL) under nitrogen at -20 °C. To the mixture were added a solution of compound 2a (142 mg, 0.132 mmol) in dry DMF (10 mL). After the mixture was stirred for 1 h at room temperature, excess trifluoroacetic acid and dichloromethane were evaporated under reduced pressure. The residue was washed with methyl-tert-butyl ether, and the supernatant was lyophilized to provide a colorless powder (109 mg). The crude yield was 72%. The product was not purified further and was used in the next reaction. MALDI-TOF-MS (matrix: α-CHCA): m/z = 571.40 [M + H]+. ¹H-NMR (CD₃CN + D₂O, δ/ppm): 3.98 (10H, s), 3.76 (10H, s), 3.5–3.7 (20H, br), 1.340 (45H, br).

N,N”,N’,N’’,N’’’,N’’’’-Pentakis(N-tetrahydroy succinimide ester (248 mg, 0.876 mmol) in acetone to provide a colorless powder. The crude product was purified by reversed-phase HPLC (column: Inertsil ODS-3) columns eluting with a linear gradient of CH₃CN/water (30/70 to 50/50 over 20 min). The isolated yield was 12%. MALDI-TOF-MS (matrix: α-CHCA): m/z = 1409.88 [M + H]+. ¹H-NMR (CD₃CN + D₂O, δ/ppm): 4.65–4.80 (5H, m), 2.9–4.2 (30H, br m), 1.2–1.3 (15H, m).
**Figure 5.** Schematic illustration of the self-assembly of trigonal and pentagonal conjugates of tryptophane zipper peptide.

![Schematic illustration of the self-assembly of trigonal and pentagonal conjugates of tryptophane zipper peptide.]

**Figure 6.** Structural change of Pentagonal-Gly-Trpzip by heating process. (a) Temperature dependence of CD spectrum for aqueous solution of Pentagonal-Gly-Trpzip (10 μM) at pH 7; (b) Temperature dependence of [θ] at 233 nm for the aqueous solution. The heating and cooling rates were 1 °C/min [41]; (c) CD spectra of Pentagonal-Gly-Trpzip at 25 °C before (green line) and after (orange line) the heating process; and (d) TEM image of Pentagonal-Gly-Trpzip at 25 °C after the heating process.

![CD spectra and TEM image of Pentagonal-Gly-Trpzip]
A carbon-coated Cu-grid (Oken Co., Ltd., Tokyo, Japan) hydrophilized for 30 sec by hydrophilizing treatment apparatus (JEOL HDT-400). The sample solutions for CD spectra were also used for TEM observation. An aliquot (10 µL) of the solutions was applied to a hydrophilized carbon-coated Cu-grid, left for 60 s, and then removed. Subsequently, a drop of 2 wt% aqueous phosphotungstic acid was placed on the grids and dried in vacuo (post-staining method). The Cu meshes were subjected to TEM observation (JEOL JEM-2010) with acceleration voltage of 120 kV at 25 °C. All of the measurements were repeated for at least two samples to ensure data reproducibility.

4. Conclusions

We have developed pentagonal conjugates of tryptophane zipper-forming peptide with a pentaazacyclotriacenedc core [Pentagonal-Gly-Trpzip and Pentagonal-Ala-Trpzip]. CD spectra revealed that Pentagonal-Gly-Trpzip formed tryptophane zipper structure at pH 7, but Pentagonal-Ala-Trpzip and the precursor peptide CKWTWTE showed the formation of weak tryptophane zipper or random coil structure. Pentagonal-Gly-Trpzip self-assembled into nanofibers with uniform width at pH 7, whereas Pentagonal-Ala-Trpzip formed irregular aggregates which reflect less formation of tryptophane zipper structure. The nanofibers from Pentagonal-Gly-Trpzip showed irreversible transformation to helical ribbon accompanying the secondary structural change by heating process. The present peptide nanostructures would be applied as biodegradable nanomaterials and platforms for nano-biotechnology. The present pentagonal molecular design would extend the feasibility of multivalent peptide conjugates.

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25. Extension of the incubation time (96 h) minimally affected the results of CD spectra and TEM images.
26. The measurement was repeated for three times to confirm data reproducibility. The standard deviation was about 2%.